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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

November 1, 2004, 03:22:05; Search time 2219 Seconds (without alignments) 18906.432 Million cell updates/sec Run on:

US-10-005-469-1 Title: Perfect score:

1 gccagccccgattgggggc......ttctctgcagatcaagtact 7992 Sequence:

4134886 segs, 2624710521 residues IDENTITY NUC Gapop 10.0 , Gapext 1.0 Scoring table: Searched:

8269772

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Total number of hits satisfying chosen parameters:

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

genesequ2000s:*
genesequ2001as:*
genesequ2001bs:*
genesequ2002bs:*
genesequ2003as:* geneseqn2003cs:* geneseqn2003ds:* N_Geneseq_23Sep04:* 1: geneseqn1980s:* geneseqn1990s:* geneseqn1980s:* Database :

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

geneseqn2004s:

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SUMMARIES		ai 	AAL47276	ARK91412	ACA61697	ADC83762	AAL47281	ABK91448	ABK91435	ABK91243	ARK91434	ANDOCACA	**************************************	ADTE: 204E	ARK91440	AD56271	AAD5333	77000	7007 144	AAD4 /2 / /	ADP86265	AAD25326	ADP86272	ABK91242	
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Novel nucleic acid encoding replication competent recombinant hepatitis C virus genome useful for screening anti-hepatitis C virus therapeutics and for vaccine development.

Claim 6; Page 43-47; 85pp; English.

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ALIGNMENTS

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Hepatitis; HCV; core-neo; NS3 proteinase/helicase; vaccine; diagnosis; virucide; hepatotropic; gene therapy; anti-viral; gene; ds.
                                                                                                                                                                                                                                                             Hepatitis C virus sub-genomic replicon clone 1377-NS3-3'UTR.
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/product= "core-neo fusion protein"
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/*tag= b
/product= "NS3 proteinase/helicase"
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342. .1181
                               AAL47276 standard; DNA; 7992 BP.
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P-PSDB; AAO18000, AAO18001.
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The present invention provides protein and coding sequences from Hepatitis C virus (HCV), comprising all or part of the HCV genome and able to replicate efficiently when transfected into a susceptible cell line without reducing the growth rate of the cell line by more than 10 fold. The sequences are useful for screening for anti-HCV therapeutics, for detecting antibodies to HCV in a biological sample such as blood, serum, plasma, blood cells, lymphocytes, or liver cells from a subject, serum, plasma, blood cells, lymphocytes, or liver cells from a subject for deriving authentic HCV components such as replication-complement non-infectious, replication-defective infection-component, and replication-defective non-infectious HCV, in gene therapy or gene vaccination defective and for treating an animal infected or susceptible to HCV infection and for studying HCV infection and propagation. The present sequence is a clone of a fragment of the HCV genome which encodes core-neo and NS3 proteinase/helicase proteins the

Sequence 7992 BP; 1648 A; 2369 C; 2243 G; 1732 T; 0 U; 0 Other;

ö ö 6; Length 7992; Indels .; 0 DB 100.0%; Score 7992; 100.0%; Pred. No. 0; cive 0; Mismatches Query Match
Best Local Similarity 100.
Matches 7992; Conservative

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	CTGGAAGACACTGAGACCCATTGACCCCCTCTGTGCCCCTCCGTGTGGAAGGGCTCTTTCCTGGAAGACCTCTGGGAAGACCTTTTCTTGTTTTTTTT	6421 GTCCAACCAGAAGAGGGGCCGCAAGCCAGCTCGCCTTATCGTATTCCCAGATTTGGGG 6480 [6481 GTTGGGGGAGAAAATGGCCCTTTACGATGTGGTCTCCACCCTCCCT	6541 AIGGGCTCTTCAIACGGATTCCAATACTCTCCTGGACAGCGGGTCGAGTTCCTGGTGAAT 6600	6601 GCCTGGAAAGCGAAGAATGCCCTATGGGCTTCGCATATGACACCCGCTGTTTTGACTCA 6660	6661 ACGGTCACTGAGAATGACATCCGTGTTGAGGAGTCAATCTACCAATGTTGTGACTTGGCC 6720	6721 CCCGAAGCCAGACAGGCCATAAGGTCGCTCACAGAGCGGCTTTACATCGGGGGCCCCTG 6780	6781 ACTAATTCTAAAGGCAGAAACTGGGGCTATCGCCGGTGCCGCGGAGCGGTGTACTGACG 6840 	6841 ACCAGCTGCGGTAATACCCTCACATGTTACTTGAAGGCCGCTGCGGCCTGTCGAGCTGC 6900	6901 AAGCTCCAGGACTGCACGATGCTCGTATGCGGAGACGACCTTGTCGTTATCTGTGAAAGC 6960 	6961 GCGGGGACCCAAGAGGACGAGCCTACAGGGCCTTCAGGAGGCTATGACTAGATAC 7020 	7021 TCTGCCCCCTGGGGACCGCCCAAACCAGAATACGACTTGGAGTTGATAACATCATGC 7080 	7081 TCCTCCAATGTGTCGCGCACGATGCATCTGGCAAAAGGGTGTACTATCTCACCGT 7140	GACCCCACCCCCTTGCGCGGGCTGCGTGCAGACAGCTGACTGTACTACCGT		7261 ACTCATTCTTCTCCATCCTTCTAGCTCAGGAACAACTTGAAAAAGCCCTAGATTGTCAG 7320 	

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NS3 or HCV NS5 encoding region, or encephalomyocarditis virus (EMCV)

internal ribosome entry site (IRES) region coding for one or more NS3, are detailed in the specification. Also included are (1) an expression are detailed in the specification. Also included are (1) an expression or excor comprising a nucleotide sequence coding for the altered nucleic cards are cards, which is transcriptionally coupled to an exogenous promoter; (2) a recombinant cell human hepatoma cell comprising the altered nucleic acids (5) a recombinant cell produced by introducing into a human hepatoma cell the altered nucleic acids; (4) producing an HCV (hepatitis C virus) cell the altered nucleic acids; (4) producing a HCV replicon enhanced cell or which containing a functional HCV replicons and HCV and HCV and host cell interactions, producing HCV RNA and cerpication enhanced cells are useful in studying HCV replication and HCV and host cell interactions, producing HCV RNA and proteins, and providing a system for measuring the ability of a compound contained may create HCV mediated diseases such as liver failure, cirrhosis and created HCV mediated diseases such as liver failure, cirrhosis and heptoma. The present sequence is the HCV based vector phycometry, used as a basis for the adaptive mutations of the invention
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             New Hepatitis C virus (HCV) replicons comprising altered HCV NS3 or HCV NS5 encoding region, or encephalomyocarditis virus (EMCV) internal ribosome entry site (IRES) region, useful in studying HCV replication and
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                                                                                              /product= "Polyprotein"
/note= "Comprising NS3, NS4A, NS4B, NS5A and NS5B"
7759. .7989
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/label= IRES
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100.0%; Score 7992;
Best Local Similarity 100.0%; Pred. No. 0;
Matches 7992; Conservative 0; Mismatches
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                Hepatitis C virus; ds; gene; thiosemicarbazone; liver inflammation;
liver failure; cirrhosis.
                                                                                                                                                                                          note= "Core-neo fusion protein selectable marker"
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/note= "HCV 3'UTR"
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                                                                                                                                                                                                                                         The invention relates to a method of treating or preventing infection by hepatitis C virus or its related conditions by delaying the onset and inhibiting replication of hepatitis C virus which comprises administering thiosemicarbazone compounds. The method is useful for treating or expectating infection by hepatitis C virus or its related conditions e.g. liver inflammation, liver failure or cirrhosis, delaying the onset and inhibiting replication of hepatitis C. The present sequence represents the hepetitis C virus expression plasmid pHCVNeol7.wt DNA
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                                                                                                                                      Treating infections by hepatitis C virus and its related conditions comprises administering thiosemicarbazone compounds.
                                                                                                                                                                                                                                                                                                                                                                                                                                  Sequence 10690 BP; 2334 A; 3045 C; 2908 G; 2403 T; 0 U; 0 Other;
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tive 0; Mismatches
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thiosemicarbazone compounds for e.g. treating and preventing is C or its related condition, and delaying the onset of hepatitis

Koch U;

Altamura S,

(ALTA/) ALTAMURA S. (KOCH/) KOCH U.

(KOCH/)

WPI; 2003-778475/73

Use of thios hepatitis C

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                                                                                                                                                                                                                          4-(cinnamyloxy) benzaldehyde thiosemicarbazone; RHEPLISA; Ia; hepatitis C; HCV replication system; bicistronic RNA replicon; neomycin phosphotransferase; human hepatoma cell line; Huh-7; neomycin pubphotransferase; human hepatoma cell line; Huh-7; neomycin sulphate; G418; pHCVNeo17.wt; replicon I377neo/NS3-3'/wt.; hepatotropic; virucide; antiinflammatory; ds.
                                                                                                                                                                          pHCVNeo17.wt plasmid containing an HCV bicistronic replicon.
                                                                                                                                                                                                                     Hepatitis C virus; thiosemicarbazone;
                                                                            BP
                                                                            ADC83762 standard; DNA; 10690
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01-JAN-2004

RESULT 4 ADC83762 301

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Synthetic. Unidentified. Hepatitis C virus.

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The invention discloses a method for the treatment and prevention of hepatitis C, or its related condition, which involves the administration of thiosemicarbazone compounds, or its salts. The inhibitory activity of of thiosemicarbazone was evaluated using selection of thiosemicarbazone was evaluated using selection is oral, parenteral (e.g. subcutaneous, intravenous, administration is oral, parenteral (e.g. subcutaneous, intravenous, intravenous, intravenous, intraversular, intrasternal injection, or infusion), by inhalation spray intraversular, intrasternal injection, or infusion), by inhalation spray intramyloxy) benzaldehyde thiosemicarbazone (Ia). The compounds cg. 4-(cinnawyloxy) benzaldehyde thiosemicarbazone (Ia). The compounds or condition, delaying the onset of hepatitis C or its related condition and inhibiting replication of the hepatitis C virus. How resplication of the hepatitis C virus. How replication of the hepatitis C virus. How replication of capable of supporting How replication can be achieved using bicistronic capable of supporting How replication can be achieved using bicistronic complication as expressing a selectable marker, the neomycin phosphotransfection of these replicons in the human hepatoma cell line, Huh-7, followed by cultivation in the presence of neomycin complication. The sequence presented is the pHCVNeol7, wt plasmid which contains the cDNA coding for an HCV bicistronic replicon identical to
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Matches 7992; Conservative
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181 GACGACCGGGTCCTTTTGGATCAACCCGCTCAATGCCTGGAGATTTGGGCGTGCCCCC 240
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              The present invention provides protein and coding sequences from Hepatitis C virus (HCV), comprising all or part of the HCV genome and able to replicate efficiently when transfected into a susceptible cell line without reducing the growth rate of the cell line by more than 10 line without reducing the growth rate of the cell line by more than 10 fold. The sequences are useful for screening for anti-HCV therapeutics, for detecting antibodies to HCV in a biological sample such as blood, serum, plasma, blood cells, lymphocytes, or liver cells from a subject, or deriving authentic HCV components such as replication-complement non-infectious, replication-defective infection-component, and replication-defective infection-component, and replication-defective infection and replication to HCV, in gene therapy or gene vaccination defective non-infectious HCV, in gene therapy or gene vaccination to HCV infection and propagation. The present sequence is a clone of a fragment of the HCV genome designated
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Novel nucleic acid encoding replication competent recombinant hepatitis C virus genome useful for screening anti-hepatitis C virus therapeutics and for vaccine development.
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Matches 7991; Conserv
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1411 CTGTIACCATACCAGGTTACGGTTACGGTCACACCTCCACCTCCACCTCTCTCT
6
2401 GCCCATCTACACCCCCTACTGGTAGCGCAAAGCACTAAAGGTCCCGCTTGCTT

QY 7861 TTTTCCTCTTTTTTTTTTTTTTTTTTTTTTTTTTTTT	RESIDENT 6 RESIDE
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The further of the following region, or encephalomy describing allered HCV internal ribosome entry site (IRES) region coding for one or more NSJ, are detailed in the specification. Also included are (1) an expression acids, which is transcriptionally coupled to an exogenous promoter; (2) a cids, which is transcriptionally coupled to an exogenous promoter; (2) a cids, which is transcriptionally coupled to an exogenous promoter; (2) a cids, which is transcriptionally coupled to an exogenous promoter; (2) a cids, which is transcriptionally coupled to an exogenous promoter; (2) a cids, which is transcriptionally coupled to an exogenous promoter; (2) a cidy in the altered nucleic acids; (3) a recombinant cell broduced by introducing into a human hepatoma cell the altered nucleic acids; (4) producing an HCV hepatitis C virus) an HCV replicon enhanced cells made in the method; and (6) measuring the ability of a compound to affect HCV activity. The HCV replicons and HCV and host cell interactions, producing HCV RNA and complication and host cell interactions, producing HCV RNA and providing a system for measuring the ability of a compound condulate one or more HCV activities e.g. to discover drugs which may hepatocellular carcinoma. The present sequence is an HCV because of vector sequence is not shown in the specification but was created by the information in Claim 16 The invention relates to nucleic acid molecules comprising altered HCV Claim 16; Page; 69pp; English

Sequence 10690 BP; 2335 A; 3045 C; 2907 G; 2403 T; 0 U; 0 Other;

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1020 1080 1080 1140 1200 1260 660 780 780 840 840 900 1260 1320 1320 1380 1440 900 960 960 1380 1500 1500 1560 1560 1620 1620 1680 1740 1680 TGCTATTGGGCGAAGTGCCGGGGCAGGATCTCCTGTCATCTCACCTTGCTCCTGCCGAGA AAGTATCCATCATGCTGATGCAATGCGGCGGCTGCATACGCTTGATCCGGCTACCTGCC CATTCGACCACCAAGGGAAACATCGCATCGAGGGAGCAGGTACTCGGATGGAAGGCGGGTC TIGICGAICAGGAIGAICIGGACGAAGAGCAICAGGGGCTCGCGCCAGCCGAACIGIICG 781 CATTCGACCACCAAGCGAAACATCGCATCGAGCGAGCACGTACTCGGATGGAAGCCGGTC CCAGGCTCAAGGCGCGCATGCCCGACGGCGAGGATCTCGTCGTGACCCATGGCGATGCCT GCTTGCCGAATATCATGGTGGAAAATGGCCGCTTTTCTGGATTCATCGACTGTGGCCGGC 961 GCTIGCCGAATATCATGGTGGAAATGGCCGCTTTTCTGGATTCATCGACTGTGGCCGGC TGGGTGTGGCGGACCGTATCAGGACATAGCGTTGGCTACCCGTGATATTGGTGAAGAGC 1021 TGGGTGTGGGGGACCCCTATCAGGACATAGCGTTGGCTACCCGTGATATTGCTGAGAGC TIGGCGGCGAAIGGGCTGACCGCTTCCTCGTGCTTTACGGTATCGCCGCTCCCGATTCGC Trescesciantesecrisacescricarescritates artescentres de la recesa de la 1141 AGCGCATCGCCTTCTATCGCCTTCTTGACGAGTTCTTGAGTTTTAAACAGACCACAACG CGAAGCCGCTTGGAATAAGGCCGGTGTGCGTTTGTCTATATGTTATTTTCCACCATATTG CCGTCTTTTGGCAATGTGAGGGCCCGGAAACCTGGCCCTGTCTTCTTGACGAGGATTCCT 1501 AACCCCCCACCTGGCGACAGGTGCCTCTGCGGCCAAAAGCCACGTGTATAAGATACACCT 1501 AACCCCCACCTGGCGACAGGTGCCTCTGCGGCCAAAAGCCACGTGTATAAGATACACCCT TGGCTCTCCTCAAGCGTATTCAACAAGGGGCTGAAGGATGCCCAGAAGGTACCCCATTGT 1621 rescricricaraceratrica caacaa caacaa caaracca a reconstration 661 781 601 721 841 901 961 1081 1021 1081 1141 1201 1201 1321 1261 1441 1561 1621 d ò ద à ð 원 ò q à g ò qq ò QQ à g à da ઠે рр g à ð 셤 g à à DP à 셤 ð 셤 ð 셤

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1681 ATGGGATCTGATCTGGGGCCTCGGTGCATTACATGTTTAGTCGAGGTTAAAA 1740			CCGAAGGGTTGCGAAGGG CCGAGGGTTTGCGAAGGG GTCCCCGGTCTTCACGG GTCCCCGGTCTTCACGG GTCCCCGGTCTTCACGG GTCCCCGGTCTTCACGG GTCCCCGGTCTTCACGG GTCTCTACACGCCCCTACGGGGT	CHIGGGRACCGGTCGGTCGCGCCCCCTAGGTTTCGGG 252 CATGGTATCGACCCTAACATCAGAACCGGGGTAAGGACCATC 258 CATGGTATCGACCCTAACATCAGAACCGGGGTAAGGACCATC 258 CATGGTATCGACCCTAACATCAGAACTGCGACGGTGGTTGC 264 ACGTACTCCACCTATGGCAAGTTTCTTGCCGACGGTGGTTGC 264 ACGTACTCCACCTATGGAAGTTTCTTGCCGACGGTGGTTGC 264 ACGTACATAATATGTGATGAGTGCCACTCAACTGACTCGACCACT 270 ATCATAATATGTGATGAGTGCCACTCAACTGACTCGACCACT 270 GTCCTGGACCCACGGAGACGGCTGGAGCTCGACTCGTCGTC 276 ACGTACATGATGTGATGAGTGCCACTCGAACTGACTCGACCACT 270 GTCCTGGACCCAAGCGGAGACGGCTGGAGCGCACTCGTCGTC 276 CTCCTGGACCCACCCACCACCCAACTCGACCTCGTCG 276 CCGGGATCGGTCACCCACCAACCAACATCGAGGGGGT 282 CCGGGATCGGTCACCCACACCAAACATCCAAGGAGGTG 282 CCGGGATCGGTCACCCACACCAAACATCCAAGGAGGTG 282 CCGGGATCGGTCACCCACACCAAACATCCAAGGAGGTG 282

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POCACCACAMA CONTRACTOR ACTIVITY OF THE PROPERTY OF THE PROPERT	7201	RESULT' / ABK91435 standard; DNA; 10690 BP. XX XX XX XX XX XX XX DT 15-NOV-2002 (first entry) XX XX XX BY Hepatitis C virus vector construct pHCVNeo.17m2. XX
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'note= "Comprising NS3, NS4A, NS4B, NS5A and NS5B"
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'product= "Core-neo fusion protein"
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note= "Plasmid derived sequences"
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WO200259321-A2.

01-AUG-2002.

16-JAN-2002; 2002WO-EP000526.

23-JAN-2001; 2001US-0263479P.

(RICE-) IST RICERCHE BIOL MOLECOLARE ANGELETTI.

De Francesco R, Migliaccio G, Paonessa G;

WPI; 2002-599793/64.

New Hepatitis C virus (HCV) replicons comprising altered HCV NS3 or HCV NS5 encoding region, or encephalomyocarditis virus (EMCV) internal ribosome entry site (IRES) region, useful in studying HCV replication and expression.

Claim 16; Page; 69pp; English.

The invention relates to nucleic acid molecules comprising altered HCV (NS3 HCV NS5 encoding region, or encephalomyocarditis virus (EMCV) internal ribosome entry site (IRES) region confor for one or more NS3, NS54, or EMCV IRES mutations, respectively. The location of the mutations of are detailed in the specification. Also included are (1) an expression acids, which is transcriptionally coupled to an exogenous promoter; (2) a cids, which is transcriptionally coupled to an exogenous promoter; (2) a recombinant cell human hepatoma cell comprising the altered nucleic acids (3) a recombinant cell produced by introducing into a human hepatoma cell the altered nucleic acids (3) a recombinant cell produced by introducing into a human hepatoma with the method and (6) measuring the replicon enhanced cells made in the method; and (6) measuring the ability of a compound to affect HCV activity. The HCV replicons and HCV expression, and HCV and host cell in studying HCV replication and providing a system for measuring the ability of a compound compound to affect HCV activity. The HCV replication and providing a system for measuring the ability of a compound compound to affect HCV activities e.g. to discover drugs which may be pression, and they activities e.g. to discover drugs which may be provided the invention. Note: The present sequence is an HCV based vector shown in the specification but was created by the indexer using the HCV x vector sequence appearing as ABR91412 and the information in Claim 16

Sequence 10690 BP; 2334 A; 3044 C; 2908 G; 2404 T; 0 U; 0 Other;

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                                                                                                                                        TAGCTGTGAAAGGTCCGTGAGCCGCTTGACTGCAGAGAGTGCTGATACTGGCCTCTGC
                                                                                                                                                                                                                                                                        HCV; ss; pHCVNeo.17m2; adaptive mutation; liver failure; cirrhosis; hepatocellular carcinoma; NS3; NS5; encephalomyocarditis virus; EMCV; internal ribosome entry site; IRES; NS5A; HCV replication; mutant.
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'note= "Comprising NS3, NS4A, NS4B, NS5A and NS5B"
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Encephalomyocarditis virus.
Escherichia coli.
Enterobacteria phage T7.
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WO200259321-A2

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The invention relates to nucleic acid molecules comprising altered HCV NS3 or HCV NS5 encoding region, or encephalomycocarditis virus (EMCV) CC NS3 or HCV NS5 encoding region, or encephalomycocarditis virus (EMCV) CC NS3, or EMCV IRES mutations, respectively. The location of the mutations vector comprising a nucleotide sequence coding for the altered nucleic coding, which is transcriptionally coupled to an exogenous promoter; (2) a cids, which is transcriptionally coupled to an exogenous promoter; (2) a recombinant cell human hepatoma cell comprising the altered nucleic acids; (3) a recombinant cell produced by introducing into a human hepatoma cell the altered nucleic acids; (4) producing an HCV (hepatitis C virus) cell the altered nucleic acids; (4) producing an HCV (hepatitis C virus) an HCV replicon enhanced cells made in the method; and (6) measuring the ability of a compound to affect HCV activity. The HCV replication and complete cells are useful in studying HCV replication and CC replicon, and HCV and host cell interactions, producing HCV RNA and CC proteins, and providing a system for measuring the ability of a compound complete one or more HCV activities e.g. to discover drugs which may complicated diseases such as liver failure, cirrhosis and providing a system for measuring the ability of a compound complete one or more HCV activities e.g. to discover drugs which may be hepatocellular carcinoma. The present sequence is an HCV become shown in the specification but was created by the indeaxer using the HCV vector seminare of the invention and the present sequence is not vector shown in the specification but was created by the indeaxer using the HCV vector.
                                                             New Hepatitis C virus (HCV) replicons comprising altered HCV NS3 or HCV NS5 encoding region, of encephalomyocarditis virus (EMCV) internal ribosome entry site (IRES) region, useful in studying HCV replication and
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Best Local Similarity 100.0%; Pred. No. 0;
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WPI; 2002-599793/64.
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RESULT 9

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New Hepatitis C virus (HCV) replicons comprising altered HCV NS3 or HCV NS5 encoding region, or encephalomyocarditis virus (EMCV) internal ribosome entry site (IRES) region, useful in studying HCV replication and
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                                                                                                  HCV; ss; pHCVNeo.17m2; adaptive mutation; liver failure; cirrhosis; hepatocellular carcinoma; NS3; NS5; encephalomyocarditis virus; EMCV; internal ribosome entry site; IRES; NS5A; HCV replication; mutant.
                                                                                                                                                                                                                                                                                                                                                                                         /product= "Polyprotein"
/note= "Comprising NS3, NS4A, NS4B, NS5A and NS5B"
replace(5337,C)
                                                                                                                                                                                                                                                                                                                                                          from ECMV"
                                                                                                                                                                                                                                                                                                                                                          site
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942. .1181
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Product= "Core-neo fusion protein"
                                                                           Hepatitis C virus vector construct pHCVNeo.17m0.
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/note= "Plasmid derived
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label= IRES
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Escherichia coli.
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cell the altered nucleic acids; (4) producing into a human hepatoma cell the altered nucleic acids; (4) producing an HCV (hepatitis C virus) replicon enhanced cell or which containing a functional HCV replicon; (5) an HCV replicon enhanced cells made in the method; and (6) measuring the ability of a compound to affect HCV activity. The HCV replicons and HCV achility of a compound to affect HCV activity. The HCV replicons and HCV replicon enhanced cells are useful in studying HCV replicons and HCV replicon and HCV and host cell interactions, producing HCV RNA and expression, and Providing a system for measuring the ability of a compound to modulate one or more HCV activities e.g. to discover drugs which may to modulate one or more HCV activities e.g. to discover drugs which may treat HCV mediated diseases such as liver failure, cirrhosis and heptove. The present sequence is an HCV based vector hepatocellular carcinoma. The present sequence is not shown in the specification but was created by the indexer using the HCV sector sequence appearing as ABK91412 and the information in Claim 16 120 9 9 recrarrides de la consecue del la consecue de la co TGCTATTGGGCGAAGTGCCGGGGCAGGATCTCCTGTCATCTCACCTTGCTCCTGCCGAGA Achencedarecertearteachecadaceaceacecececrareineerede CGACGGGCGTTCCTTGCGCAGCTGTGTCGACTTGTCACTGAAGCGGAAGGGACTGGC GCGAGACTGCTAGCCGAGTAGTTGGGTCGCGAAAGGCCTTGTGGTACTGCCTGATAGG CGGCCGCTTCGGAGAGGGTATTCGGCTATGACTGGGCACAACAGACAATCGGCTGCT CTGATGCCGCCGTGTTCCGGCTGTCAGCGCAGGGGCGCCCGGGTTCTTTTGTCAAGACCG crdardececererrecederereadececadedececececerrerrirateaadadee GCGAGACTGCTAGCCGAGTAGTGTTGGGTCGCGAAAGGCCTTGTGGGTACTGCCTGATAGG GTGCTTGCGAGTGCCCCGGGAGGTCTCGTAGACCGTGCACCATGAGCACGAATCCTAAAC CTCAAAGAAAAACCAAAGGGCGCGCGTTGATTGAACAAGATGGATTGCACGCAGGTTCTC TCTTCACGCAGAAAGCGTCTAGCCATGGCGTTAGTATGAGTGTCGTGCAGCCTCCAGGAC CCCCCTCCCCGGGAGAGCCATAGTGGTCTGCGGAACCGGTGAGTACACCGGAATTGCCAG GACGACGGGTCCTTCTTGGATCAACCGCTCAATGCCTGGAGATTTGGGCGTGCCCCC GCCAGCCCCCGATTGGGGGGGACACTCCACATGATCACTCCCCTGTGAGGAACTACTG Gaps DB 6; Length 10690; Sequence 10690 BP; 2335 A; 3044 C; 2908 G; 2403 T; 0 U; 0 Other; 0; 1; Indels Score 7990.4; Pred. No. 0; 0; Mismatches Query Match 100.0%; Best Local Similarity 100.0%; Matches 7991; Conservative 0. 601 661 721 361 421 481 481 541 601 301 361 421 181 241 241 301 61 61 121 121 181 g ò à g ð qq à 엄 ò g ð g 8 8 8 Д ò 상 업 qq ð

1861 AGCCTCACAGGCCGGGACAGCAACCAGGTCGAGGGGAGGTCCCAAGTGGTCTCCACCGCA 1920 GCAAAGGCGGCACAACCCCCAGTGCCACGTTGTGGATAGTTGTGGAAAAGAGTCAAA 1620 AACGTCTAGGCCCCCGGAACCACGGGGACGTGGTTTTCCTTTGAAAAACACGATAATACC AIGGCGCCIATTACGGCCTACTCCCCAACAGACGCGAGGCCTACTIGGCTGCATCATCAC TGGCTCTCCTCAAGCGTATTCAACAAGGGCTGAAGGATGCCCAGAAGGTACCCCATTGT AACCCCCCACCAGGGGACAGGTGCCTCTGCGGGCCAAAGCCACGTGTATAAGATACACCT GCAAAGGCGGCACAACCCCAGTGCCAACGTTGTGAGTTGGATAGTTGTGGAAAGAGTCAAA **ATGGGATCTGATCTGGGGCCTCGGTGCACATGCTTTACATGTGTTTAGTCGAGGTTAAAA** CGAAGCCGCTTGGAATAAGGCCGGTGTGCGTTTGTCTATATGTTATTTCCACCATATTG CCGICITITIGGCAATG1GAGGGCCCGGAAACC1GGCCCTGTCTTCTTGACGAGCATTCCT AGCGCATCGCCTTCTATCGCCTTCTTGACGAGTTCTTCTGAGTTTAAACAGACCACAACG ĠĊŢŢĠĊĊĠĂĀŢĀŢĊĀŢĠĠŢĠĠĀĀĀĀŢĠĠĊĊĠĊŤŢŢŢĊŢĠĠĀŢŢĊĀŢĊĠĊŢĠĠĊĊĠĠĊ GCTTGCCGAATATCATGGTGGAAAATGGCCGCTTTTCTGGATTCATCGACTGTGGCCGGC TGGGTGTGGCGGACCGCTATCAGGACATAGCGTTGGCTACCCGTGATATTGCTGAAGAGC CCAGGCTCAAGGCGCGCATGCCCGACGCGAGGATCTCGTCGTGACCCATGGCGATGCCT CATTCGACCACCAAGGAAACATCGCATCGAGCGAGCACGTACTCGGATGGAAGCCGGTC TIGICGAICAGGAIGAACGAAGGAGCAICAGGGGCICGCGCCAGCCGAACIGIICG 1741 1801 1801 1501 1561 1621 1621 1681 1741 1501 1561 1441 1321 1081 1141 1141 1201 1201 1261 1261 1321 1381 1021 1081 196 841 901 901 961 781 781 qq ð qq g Dp δ g ð Db ₽ g à ò g δ ò g à g ₽ Q ∂ g δ 셤 à g ò g à g à 셤 8 8 ₹ 720 720 780 780 099 9 420 480 480 540 540 600 900 360 360 420 240 240 300

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		2281 ACCCGAGGGTTGCGAAGCGTGGACTTTGTACCGTCGAGTCTATGGAACCACTATG 2341 CGGTCCCCGGTCTTCACGGACAACTCGTCCCCTCCGGCCGTACCGCAGACATTCCAGGTG 2340 2341 CGGTCCCCGGTCTTCACGGACAACTCGTCCCCTCCGGCCGTACCGCAGACATTCCAGGTG 2400 2401 GCCCATCTACACGACAACTCGTCCCCTCCGGCCGTACCGCGGCTATGCA 2460 2401 GCCCATCTACACGCCCTACTGGTAGCGCAAGAGCACTAAGGTGCCGGCTATGCA 2460 2401 GCCCATCTACACGCCCTACTGGTAGCGCAAGAGCACTAAGGTGCCGGCTATGCA 2460 2461 GCCCAAGGGTATAAGGTGCTTGTCCTGAAGCCGCACCCCTAAGGTTTCGGG 2520 2461 GCCCAAGGGTATAAGGTGCTTGTCCTGAACCCGTCGCCCCCCACGCTATGCA 2460 2461 GCCCAAGGGTATAAGGTGCTTGTCCTGAACCCGTCGCCCCCCCACAGGTTTCGGG 2520	GCGTATATGTCTAAGGCACATGGTATCGACCCTACATCGCCGCCACCCTAGGTTTCGGG GCGTATATGTCTAAGGCACATGGTATCGACCCTAACATCAGAACCGGGGTAAGGATATCGGC GCGTATATGTCTAAGGCACATGGTATCGACCCTAACATCAGAACCGGGGGTAAGGACCATC ACCACGGGTGCCCCCATCGTATGGCCAACTTTCTTGCCGACGGTGGTTGC	2701 ATCCTGGGCATCGGCACAGTCCTGGACGAGGGGAGGCGGGCG	294

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4.081 GCGATTAGCATCACTGATGGCATTCACAGCCTCTATACCAGCCCGCTCTACCCCCCCC

7321 7381 7381 7441 7441 7501	0.9 7561 AACTGGGCACTAAGGACCAAGCTCACTCCCAATCCCGGCTGCGTTGGAT 7620 1		7921 7921 7981 7981	986264 986264 ADP86264; ADP86264; 23-SEP-2004 (first entity) Hepatitis C virus Con-1	<pre>KW Hepatitis C virus; HCV; anti-HCV agent; HCV infection; therapy; plasmid; XX XX XX</pre>
	6481 GTTCGTGTGTGCGAGAAATGGCCCTTTACGATGTGGTCTCCACCCTCCAGGCGGG 6540 [9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9		6961 GCGGGGACCCAAGAGGACGAGCCTACGGGCCTTCACGGAGGCTATGACTAGATAC 7020 [AAT

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The present invention provides hepatitis C virus (HCV) replication cells and cell lines derived from human non-hepatic cells or non-human cells. The invention is useful for identifying anti-HCV agents for treating HCV infections. The present sequence is hepatitis C virus Con-1 replicon 1377/NS3-3' plasmid DNA.
                            at replicates hepatitis C virus (HCV), where the cell from a non-human cell line and a human non-hepatic cell identifying anti-HCV agents for treating HCV infections.
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ilarity 100.0%; Pred. No. 0;
Conservative 0; Mismatches
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1861 AGCCTCACAGGCCGGGACAGGACCAGGTCGAGGGCCCAAGTGGTCCCCACCGCA 1920		GGCGGTCCACTGCTCTGCCCCTCGGGGCACGCTGTCTCTACTTGAAGGGCTGGCGGCTCTCCTACTTTGAAGGGCTGGCGGCTGCGGGCTGGGGCTGTGGGCCTGTGTGGGCTTGTGGGCCTGCGGGCACGCTGTGGGGCATCTTTCGGGCCTGCGGGCACGCTGTGGGGCATCTTTTGGGGCTTGTGGGCTTGTGTGGGCTTGTGGGAAACCGAGGGGTTGGTGGACTTTGTACCGAGGGGTTGTATGGAAACCAACGAGGGGTTGGTGGGACTTTGTACCGAGGGGTTGTATGGAAACCAACGAGGGGTTGGTGGACTTTGTACCGAGGGGTTGTATGGAAACCAACAAGGGGGTTGGTT	AGGTG AGGTG ATGCA	GCCCAAGGGTATAAGGTGCTTGTCCTGAACCCGTCGCGCCGCCGCCTGCGCTGCCCGCCTGCGCTGCCCGCCACCCTAGGTT	311GC 26 311GC 26 31TGC 26	ATCCTGGGCATCGGCACAGTCCTGGACCAGCGAGACGCCTGAGCTCGACCCCT	GCTCTGTCCAGCACTGGAGAAATCCCCTTTTATGGCAAAGCCATCCCATCGAGAGACCCTGTCCAGGAGAAGCCATCCCCTTTTATGGCAAAGCCATCCCCATCGAGAGACTCCCTTTTATGGCAAAGCCATCCCATCGAGAAGCCATCCCATCGAGAAAGCCATCCCATCGAGAAAGCCATCCCATCGAGAAAGCCATCCCATCGAGAAAGCCATCCCATCGAGAAAGCCATCCCATCGAGAAAGCCATCCCATCGAGAAAGAA	CCGCG 294	2941 AAGCTGTCCGGCCTCGGACTCAATGCTGTAGCATATTACCGGGGCCTTGATGTATCCGTC 3000

ACAGACCCCTGATCACGCCATGCGCTGCGGAGGAACCAAGCTGCCCATCAATGCACTG AGCAACTCTTTGCTCCGTCACCACAACTTGGTCTATGCTACAACATCTCGCAGCGCAAGC CCCATATGGGCACGCCGGATTACAACCCTCCACTGTAGAGTCCTGGAAGGACCCGGACCCGGACCCGGACCCCGGACCCCGGACCCCGGACCCTCCACTTAGAGTCCTGGAAGGACCCGGACCCGGACCCGGACCCGGACCCGGACCCGGACCCGGAACGACCCTCCACTTAGAGTCCTGGAAGGACCCGGAC ACGCCTCTCCTGACCACCCTCCGACGGCGACGCGGGGATCCGACGTTGACTCGTAC CTGCGGCAGAAGAAGGTCACCTTTGACAGACTGCAGGTCCTGGACGACCACTACCGGGAC AGGGAAGTATCCGTTCCGGCGGAGATCCTGCGGAGGGACCCAGGAATTCCCTCGAGCGATG TACGTCCTCCAGTGGTACACGGGTGTCCATTGCCGCCTGCCAAGGCCCCTCCGATACCA GCTAAGCGTAGGCTGGCCAGGGATCTCCCCCTCCTTGGCCAGCTCATCAGCTAGCCAGGCTAGCCAGGCTAGCCAGGCTAGCCAGGCTAGCCAGGGATCTCCCCCCTCCTTGGCCCAGCTATCAGCTAGCCAGGCAAGAATCTCCCCCCTCCTTGGCCAGCTATCAGCTAGCCAG CTGTCTGCGCCTTCCTTGAAGGCAACATGCACTACCCGTCATGACTCCCCGGACGCTGAC CTCATCGAGGCCAACCTCCTGTGGCGGCAGGAGGATGGGCGGGAACATCACCCGGCGTGGAG ccesacerraceagrecreacricearecreacecrecerecacacracacres 5161 ACATTCCTGGTCGGGCTCAATCATACCTGGTTGGGTCACAGCTCCCATGCGAGCCCGAA 원 수 명 П 印格 军格 甲格 甲格 甲格 甲格 甲格 甲格 甲 6 8 6 8 6 8 6 ò

131	RESULT 11 AAA98968 1D AAA98968 standard; DNA; 7989 BP. XX AC AAA98968; XX DT 08-FEB-2001 (first entry) XX DE Hepatitis C virus DNA fragment SEQ ID NO: 4. XX
6241 GAAGCCTGTAAGCTGACGCCCCACATTCGGCCAGATCTAAATTTGGCTATGGGCCAAG 6300	6961 GCGGGGACCCAAGAGGACGAGCCTACGGGCCTTCACGGAGGCTATGACTAGATAC 7020 6961 GCGGGGACCCAAGAGGCGAGCCTACGGGCCTTCACGGAGGCTATGACTAGATAC 7020 6961 GCGGGGACCCAAGAGGCCTACGGGCCTTCACGGGGCTATGACTACATAC 7020 7021 TCTGCCCCCCTGGGGACCCGCCCAAACCAATACGACTTGAGATAACATCATGC 7080 7021 TCTGCCCCCCTGGGGGACCCGCCCAAACCAATACGACTTGAGATAACATCATGC 7080 7081 TCTCCCCAATGTGACGCGCCCCAAACCAAAAGGCTATAAACATCACCCG 7140 7081 TCCTCCAATGTGACGCGCCCCCAAACCAAAAGGCTAACTATCTCACCCGT 7140 7081 TCCTCCAATGTGACGCGCCCCAAACCAAAAGGCTAACTATCTCACCCGT 7140 7141 GACCCCACCACCCCCTTGCGCGCGCGCGCGCGAAAAGGCTAACTATCTCACCCGT 7140 7141 GACCCCACCACCCCCCTTGCGCGCGCGCGCGCGCAAAAGGCTAGACAACTCCAGTCAAT 7200 7141 GACCCCACCACCCCCCTTGCGCGCGCGCGCGCGCGAAAGGCTAGACACCCCAGTCAAT 7200 7201 TCCTGGCTAGGCAACATCATGTATGCGCCCACCTTGTGGGCAAGGATGATCCTGATG 7260 7201 TCCTGGCTAGGCAACATCATGTATGCGCCCACCTTGTGAAAAAGCCCTAGATTGTCAAGAGGTGATCTTGTAGAAAAAGCCTTGAAAAAAGCCTTAGAAAAAAGCCTTAGAAAAAAGCCTTAGAAAAAAGCCTTAGAAAAAAGCCTTAGAAAAAAGCCTTAGAAAAAAGCCTTAGAAAAAAAGCCTTAGAAAAAAGCCCTAGATTGTCAACTCCAATCTATGAAAAAAACCCTTAGAAAAAAACCCTTAGAAAAAAACCCTTAGAAAAAAACCCTTAGAAAAAAACCCCTTAGAAAAAAACCCTTAGAAAAAAACCCTTAGAAAAAAACCCTTAGAAAAAAACCCTTAGAAAAAAACCCTTAGAAAAAAAA

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Cell culture system for hepatitis C virus, useful e.g. in screening for therapeutic agents, comprises human hepatoma cells containing a viral RNA construct that includes a selectable gene.

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8; Page 37-43; 58pp; German.

This invention describes a novel Hepatitis C virus (HCV) cell culture system comprising human hepatoma cells that contain an integrated HCV-RNA construct [1]. (I) contains the HCV-specific RNA segments 5. WTR (non-translated region), NS (non-structural)3, NS4B, NS4B, NS4B, NS5B and 3. NTR, and a selectable (marker) gene [II). The cell cultures, and/or [I). The cell cultures and/or [I). The cell cultures and/or test therapeutic and/or diagnostic agents for HCV infections, and to prepare vaccines against HCV infection (particularly preparation of attenuated HCV). The can also be used for preparation of a liver-specific delivery system for gene therapy, and to preparation of a termaneth HCV). The can also be used for preparation of a liver-specific delivery system for gene therapy, and to identify cells permissive for HCV replication. Virus rystem, so that autonomously and with high efficiency in this cellular system, so that variations in replication rates can be measured (for screening antiviral agentis) quantitatively, using standard laboratory equipment. Efficient replication of HCV RNA is only achieved when the specified RNA segments are present and when the transfected cells are maintained under permanent selection pressure

Sequence 7989 BP; 1647 A; 2368 C; 2243 G; 1731 T; 0 U; 0 Other;

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480 540 009 360 GTGCTTGCGAGTGCCCCGGGAGGTCTCGTAGACCGTGCACCATGAGCACGAATCCTAAAC 360 420 420 480 Accigiocogogocorda Arga Accide a Gorda Gor CGACGGGGGTTCCTTGCGCAGCTGTGCTCGACGTTGTCACTGAAGGGGAAGGGACTGGC 660 TCTTCACGCAGAAAGCGTCTAGCCATGGCGTTAGTATGAGTGTCGTGCAGCCTCCAGGAC 120 GCGAGACTGCTAGCCGAGTAGTGTTGGGTCGCGAAAGGCCTTGTGGTACTGCCTGATAGG 300 9 CTGATGCCGCCGTGTTCCGGCTGTCAGCGCAGGGGCGCCCGGTTCTTTTGTCAAGACCG GCCAGCCCCCGATTGGGGGGGACACACCACCATAGATCACTCCCCTGTGAGGAACTACTG TCTTCACGCAGAAAGCGTCTAGCCATGGCGTTAGTATGAGTGTCGTGCAGCCTCCAGGAC CCCCCCTCCCGGGAGAGCCATAGTGGTCTGCGGAACCGGTGAGTACACCGGAATTGCCAG CCCCCTCCCGGGAGAGCCATAGTGGTCTGCGGAACCGGTGAGTACACCGGAATTGCCAG GCCAGCCCCCGATTGGGGGGGACACTCCACCATAGATCACTCCCCTGTGAGGAACTACTG GACGACCGGGTCCTTTCTTGGATCAACCCGCTCAATGCCTGGAGATTTGGGCGTGCCCCC GACGACCGGGTCCTTTCTTGGATCAACCCGCTCAATGCCTGGAGATTTGGGCGTGCCCCC GCGAGACTGCTAGCCGAGTAGTGTTGGGTCGCGAAAGGCCTTTGTGGTACTGCCTGATAGG GTGCTTGCGAGTGCCCCGGGAGGTCTCGTAGACCGTGCACCATGAGCACGAATCCTAAAAC CTCAAAGAAAAACCAAAGGGCGCGCCATGATTGAACAAGATGGATTGCACGCAGGTTCTC CTCAAAGAAAACCAAAGGGGGGGCGCATGATTGAACAAGATGGATTGCACGCAGGTTCTC CGGCCGCTTGGGTGGAGGCTATTCGGCTATGACTGGGCACAACAGACAATCGGCTGCT CGCCCGCTTGGGTGGAGAGGCTATTCGGCTATGACTGGGCACAACAACAGACAATTCGGCTGCT ô Length 7989; 0; Indels 3; DB Pred. No. 0; 0; Mismatches 100.0%; Score 7989; 100.0%; Pred. No. 0; Best_Local Similarity 100. Matches 7989; Conservative 241 541 541 601 Query Match 61 61 121 121 181 181 241 301 301 361 361 421 421 481 481

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Qy 7201 TCCTGGCTAGGCAACATCATCATGTATGCGCCCACCTTGTGGGCAAGGATGATCCTGATG 7260 Db 7201 TCCTGGCTAGGCAACATCATGTATGCGCCCACCTTGTGGGCAAGGATGATCCTGATG 7260 Qy 7261 ACTCATTCTTCTCCATCCTTCTAGGTCAGGAACAACTTGAAAAAGCCCTAGATTGTCAG 7320 Db 7261 ACTCATTCTTCTCCATCCTTCTAGGTCAGGAACAACTTGAAAAAGCCCTAGATTGTCAG 7320 Qy 7321 ATCTACGGGGCCTGTTAGTCCATTGAGCTACCTCAGATTGTCAG 7320	7321 ATCTACGGGGCTGTTACTCCATAGAGCCACTTGACCTACCT	Db 7441 TCATGCCTCAGGAAACTTGGGGTACCGCCCTTGCGAGTCTGGGGCCAGAAGT 7500	7621 TTATCCAGCTGGTTCGTTGCTGGTTACAGCGGGGGGGACATATATCACAGCCTGTCTGT	7741 7741 7801		7981 AGATCAAGT 7989 7981 AGATCAAGT 7989 SULT 12	XX XA ADJ57845; Standard; DNA; 7989 BP. XX AC ADJ57845; XX XX XX DT 06-MAY-2004 (first entry) XX XX XX DE HCV replicon encoding sequence. XX
	6241 GAAGCCTGTAAGCTGACGCCCCACATTCGGCCAGATCTAAATTTGGCTATGGGCAAAG 6300 6241 GAAGCCTGTAAGCTGACGCCCCACATTCGGCCAGATCTGACTATGGGCAAAG 6300 6241 GAAGCCTGTAAGCTGACGCCCCCACATTCGGCCAGATCTGGGTATGGGCAAAG 6300 6301 GACGTCCGGAACCTATCCAGCAAGGCGTTAACCACATCCGCTCCGTGTGGAAGGACTTG 6360 6301 GACGTCCGGAACCTATCCAGCAAGGCGTTAACCACATCGGCTCGTGGAAGGACTTG 6360 6301 GACGTCCGGAACCTATCAGCAATTGACCACTTAACCACATCGGCAAGGACTTG 6360 6361 CTGGAAGACACTGAGACCAATTGACACCACCATCATGGCAAAAAATGAGGTTTTCTGC 6420	6361 CTGGRAGACACCAATTGACCCACCACCAGGCAAAAAATGAGGTTTTCTGC 6420 6421 GTCCAACCAGAGAGAGGGGGCGCAAGCCACCATTATCGTATTCCCAGATTTGGGG 6480 6421 GTCCAACCAGAGAGGGGGCGCAAGCCAGCTTGTCGTATTCCAGATTTGGGG 6480 6421 GTCCAACCAGAGAGAGGGGGCGCAAGCCAGCTTATCGTATTCCCAGATTTGGGG 6480 6481 GTTCGTGTTGTGCGGAAAATGGCCCTTTACGATGTGGCTCTCCACCCTCCAGGCGTG 6540 6481 GTTCGTGTTGTGCGAAAAATGGCCCTTTACGATGTGGTCTCCACCCTCCAGGCGTG 6540 6481 GTTCGTGTTGTGCGAAAAATGGCCCTTTACGATGTGGTCTCCACCCTCCAGGCGTG 6540		6661 ACGGTCACTGAGAATGACATCCGTGTTGAGGAGTCAATGTTGTGATGTTGGCC	6781 ACTAATTCTAAAGGGCAGAACTGCGGCTATCGCCGCGCGCG	01 AAGCTCCAGGACTGCACGATGCTCGTATGCGGAGACGACCTTGTCGTTATCTGTGAAAGC 696	021 021 081 081 141

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Best Local Similarity 100.0%; Pred. No. 0;
Matches 7989; Conservative 0; Mismatches
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  Location/Qualifiers
1801. .7759
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                                                              /product= "HCV
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The invention relates to nucleic acid molecules comprising altered HCV NS3 or HCV NS5 encoding region, or encephalomyocarditis virus (EMCV) internal ribosome entry site (IRES) region coding for one or more NS3, NS5A, or EMCV IRES mutations, respectively. The location of the mutations
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            New Hepatitis C virus (HCV) replicons comprising altered HCV NS3 or HCV NS5 encoding region, or encephalomyocarditis virus (EMCV) internal ribosome entry site (IRES) region, useful in studying HCV replication and
                                                                                                                 HCV; ss; pHCVNeo.17m2; adaptive mutation; liver failure; cirrhosis; hepatocellular carcinoma; NS3; NS5; encephalomyocarditis virus; EMCV; internal ribosome entry site; IRES; NS5A; HCV replication; mutant.
                                                                                                                                                                                                                                                                                                                                                                                              /note= "Comprising NS3, NS4A, NS4B, NS5A and NS5B"
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/label= IRES
tote= "Internal ribosome entry site from ECMV"
1801. .7758
                                                                                                                                                                                                                                                                                               "Core-neo fusion protein"
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                        ABK91440 standard; DNA; 10690
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Encephalomyocarditis virus.
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Enterobacteria phage T7.
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5'UTR
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care detailed in the specification. Also included are (1) an expression vector comprising a nucleotide sequence coding for the altered nucleic acids acidis, which is transcriptionally coupled to an exogenous promoter; (2) a recombinant cell human hepatoma cell comprising the altered nucleic acids; (3) a recombinant cell produced by introducing into a human hepatoma cell comprising an HCV (hepatitis C virus) replicon enhanced cells and entitly of a compound to affect HCV activity. The HCV replicons and HCV replicon enhanced cells are useful in studying HCV replicons and HCV replicons and HCV replicons, and HCV and host cell interactions, producing HCV RNA and expression, and HCV and host cell interactions, producing HCV RNA and providing a system for measuring the ability of a compound complete one or more HCV activities e.g. to discover drugs which may treat HCV mediated diseases such as liver failure, cirrhosis and hepatoma carcinoma. The present sequence is an HCV based vector pHCVNeo.17 mutant of the invention. Note: The present sequence is not shown in the specification but was created by the indexer using the HCV vector sequence appearing as ABK91412 and the information in Claim 16 540 240 300 300 360 480 009 720 ó. 180 240 360 420 420 480 540 600 099 099 720 cececerecessas de contra de contra de cecesa de contra de cece de contra de cecesa de contra de cecesa de contra de cecesa de 120 TCTTCACGCAGAAAGCGTCTAGCCATGGCGTTAGTATGAGTGTCGTGCAGCCTCCAGGAC 120 09 9 CCCCCCTCCCGGGAGAGCCATAGTGGTCTGCGGAACCGGTGAGTACACCGGAATTGCCAG GCGAGACTGCTAGCCGAGTAGTGTTGGGTCGCGAAAGGCCTTGTGGTACTGCCTGATAGG GTGCTTGCGAGTGCCCCGGGAGGTCTCGTAGACCGTGCACCATGAGCACGAATCCTAAAAC CTGATGCCGCCGTGTTCCGGCTGTCAGCGCAGGGCCGCCCGGTTCTTTTTGTCAAGACCG creareccecererreceecrercaececaececececeerrerrrrrercaaeaece CGACGGGCGTTGCGCTGTGCTCGACGTTGTCACTGAAGCGGAAGGGACTGGC TGCTATTGGGCGAAAGTGCCGGGGCAGGATCTCCTGTCATCTCACCTTGCTCCTGCCGAGA GCCAGCCCCCGATTGGGGGCGACACTCCACCATAGATCACTCCCCTGTGAGGAACTACTG GACGACCGGGTCCTTTCTTGGATCAACCCGCTCAATGCCTGGAGATTTGGGCCGTGCCCCC GTGCTTGCGAGTGCCCCGGGAGGTCTCGTAGACCGTGCACCACCATGAGCACGAATCCTAAAC CTCAAAGAAAAACCAAAGGGCGCGCCATGATTGAACAAGATGGATTGCACGCAGGTTCTC CGGCCGCTTGGGTGGAGGCTATTCGGCTATGACTGGGCACAACAGACAATCGGCTGCT 541 Accrercedrecerrehardaacrecadaadededecedecederrarderegeera 1 GCCAGCCCCCGATTGGGGGGGGCACTCCATCATAGATCACTCCCCTGTGAGGAACTACTG TCTTCACGCAGAAAGCGTCTAGCCATGGCGTTAGTATGAGTGTCGTGCAGCCTCCAGGAC DB 6; Length 10690; Sequence 10690 BP; 2333 A; 3045 C; 2908 G; 2404 T; 0 U; 0 Other; ; 0 2; Indels 100.0%; Score 7988.8; ilarity 100.0%; Pred. No. 0; Conservative 0; Mismatches Best Local Similarity Matches 7990; Conserv 661 61 61 121 121 181 181 241 301 301 361 361 421 421 481 481 601 501 661 Query Match g g g d ò g $\dot{\delta}$ 셤 ð qq à à g à 8 à qq à ò à 유 ð g à

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                                                                                                                        The present invention provides hepatitis C virus (HCV) replication cells and cell lines derived from human non-hepatic cells or non-human cells. The invention is useful for identifying anti-HCV agents for treating HCV infections. The present sequence is hepatitis C virus Con-1 replicon 1377/NS3-3' derived plasmid DNA.
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                                                               New cell-line that replicates hepatitis C virus (HCV), where the cell line is selected from a non-human cell line and a human non-hepatic cell line, useful for identifying anti-HCV agents for treating HCV infections.
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/product= "HCVreplbBartMan polyprotein"
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The invention relates to Hepatitis C virus (HCV) variants which include variants that have a transfection efficiency and ability to survive contribute that have a transfection efficiency and ability to survive subpassage greater than HCV that have wild-type polyprotein coding regions. The polymucleotides of the invention are useful for identifying a call line that is permissive for infection with HCV and detecting replication of HCV in cells of the cell line. They are also useful for replication. They are also useful for the sting a compound for anti-viral properties and for inhibiting HCV infection. They are also useful for the generation of defined HCV virus crocks to develop in vitro and in vivo assays for virus neutralisation, controlled and RNA elements and identification of new antiviral targets, proteins and RNA elements and identification of new antiviral targets, those that support wild-type and variant HCV RNA replication and particle treass, production of adaptive HCV variants capable of more efficiency replication in cell culture, production of alternative animal models for inhibitor evaluation including those supporting HCV variant replication, cinhibitor evaluation including those supporting HCV variant replication, development of cell-free HCV replication, assays, production of alternated HCV immunospanic HCV particles for vaccination, engineering of attenuated HCV derivatives as noweithe vaccination, environment of cell-free HCV replication, assays, production of adaptive HCV received and particles for vaccination, engineering of attenuated HCV derivatives as noweithe vaccination, environment of cell-free HCV replication, and particles for accination, engineering of attenuated HCV derivatives as noweithe vaccinations environment and particles for accination, environment of cell-free HCV received environment o
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Hepatitis C virus variants having greater transfection efficiency and ability to survive subpassage, useful as a vaccine for immunizing primate to the virus, comprise non-naturally occurring viral sequences.
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Search completed: November 1, 2004, 19:16:14

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Hepatitis C virus replicons and replicon enhanced cells
Patent: WO 02059121-A 3 01-AUG-2002;
ISTITUTO DI RICERCHE DI BIOLOGIA MOLECOLARE P. ANGELETTI (IT)
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(c) 1993 - 2004 Compugen Ltd.
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	3721 GTCCTAGCAGCTCTGGCGGTATTGCCTGACAACAGGCGGGGTCATTGTGGGGCAGG 3780

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REFERENCE 1 AUTHORS Bartenschlager,R.D. TITLE Hepatitis C virus cell culture system JOURNAL Patent: EP 1043399-A 4 11-OCT-2000; FEATURES Location/Qualifiers Source /organism="Hepatitis C virus" /mol_type="unassigned DNA" ORIGIN AUD_XTEE="Laxon:11103"	Query Match 100.0%; Score 7989; DB 6; Length 7989; Best Local Similarity 100.0%; Pred. No. 0; Matches 7989; Conservative 0; Mismatches 0; Indels 0; Gaps 0; QY 1 GCCAGCCCCGATTGGGGGCGACACTCCACCATAGATCACTCCCCTGTGAGGAACTACTG 60 Indept. Ind	Db		181	241	OY 301 GTGCTTGCGAGTGCCCGGGAGGTCTCGTAGACCGTGCACCATGAGCACGAATCCTAAAC 360 Db 301 GTGCTTGCGAGTGCCCCGGGAGGTCTCGTAGACCGTGCACCATGAGCACGAATCCTAAAC 360	QY 361 CTCAAAGAAAAACCAAAGGGCGCGCCATGATTGAACAAGATGGATTGCACGCAGGTTCTC 420	Qy 421 CGGCCGCTTGGGTGGAGAGCTATTCGGCTATGACTGGCACAACAACAACAGCTGCT 480 L	Oy 481 CTGATGCCGCCGTGTTCCGGCTGTCAGGGGCGCCCGGTTCTTTTGTCAAGACG 540	QY 541 ACCTGTCGGTGCCCTGAATGAACTGCAGGGCAGCGGGGCTATCGTGGCTGG 600 Db 541 ACCTGTCCGGTGCCCTGAATGAACTGCAGGACGAGGGGGGTATCGTGGCTG 600	OY 601 CGACGGGCTTCCTTQCGCAGCTGTQCTCGACTGAAGCGGAAGGACTGGC 660	661	Oy 721 AAGTATCCATCATGGCTGATGCAATGCGGCGGCTGCATACGCTTGATCCGGCTACCTGCC 721 AAGTATCCATCATGATGCAATGCGGCGGCTGCATACGCTTGATCCGGCTACCTGCTTGATCCGGCTACCTGCC 780	Oy 781 CATTGACCACCAAGGAAACATGCATCGAGGGAGCAGGTACTGGAAGGCGGTC 840	Qy 841 TTGTCGATCAGGATGATCTGGACGAGCATCAGGGGCTCGCGCCAGCCGAACTGTTCG 900
Qy 7141 GACCCCACCACCACCACCCCCTTGGGGGGTGGGGAGACAGCTAGACCACTCCAGTCAAT 7200 Db 7141 GACCCCACCACCCCCTTGGGGGGTGGGGGGAGACAGCTAGACACTCCAGTCAAT 7200 Qy 7201 TCCTGGCTAGGCAACATCATCATGTATGCGCCCACCTTGTGGGCAAGGATGATCCTGATG 7260 Db 7201 TCCTGGCTAGGCAACATCATGTATGCGCCCACCTTGTGGGCAAGGATGATCCTGATG 7260 Qy 7261 ACTCATTCTTCTCCATCCTTCAGCTCAGGAACAACTTGAAAAAGCCCTAGATTGTCAG 7320 Db 7261 ACTCATTCTTCTCCATCCTTCAGCTCAGGAACAACTTGAAAAAAGCCCTAGATTGTCAG 7320 Db 7261 ACTCATTCTTCTCCATCCTTCTAGCTTCAGAAAAAGCCCTAGATTGTCAG 7300	738	Db 7381	7501 GTCGCGCTAGGCTACTGTCCCAGGGGGGGGGGGGGTGCCACTTGTGGCAAGTACCTCTTC 7	VY 7561 AACTGGGCAGTAAGGACCAAGCTCAACTCCAATCCCGGCTGCGTCCCAGTTGGAT 7620	OY 7621 TTATCCAGCTGGTTGCTGGTTACAGCGGGGGAGACATATATCACAGCCTGTCTCGT 7680	QY 7681 GCCCGACCCGCTGGTTCALGTGCCTACTCCTACTTTCTGTAGGGTAGG	Qy 7741 CTACTCCCCAACGATGAACGGGGAGCTAAACACTCCAGGCCAATAGGCCATCCTGTTTT 7800	OY 7801 TITCCTITITITITITITITITITITITITITITITITI	QY 7861 TITITCCTTITITCTTITCTTITCTTITGGIGGCTCCARCTIAGCCCTAGTCACGC 7920	OY 7921 IAGCTGTGAAAGGTCCGTGAGCCGCTTGACTGCAGAGTGCTGATACTGGCTCTCTGC 7980		NV03C3E	DEFINITION Sequence 4 from Patent EP1043399. ACCESSION AX036255.1 GI:11225871	Hepa Hepa Viru	

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TCCTGGCTAGGCAACATCATCATGTATGCGCCCACCTTGTGGGCAAGGATGATCCTGATG ACCAGCTGCGGTAATACCCTCACATGTTACTTGAAGGCCGCTGCGGCCTGTCGAGCTGCG GCGGGGACCCAAGAGGACGAGCCGACCTTCACGGGGCTTATGACTAGATAC GACCCCACCACCCCCTTGCGCGGGCTGCGTGGGAGACAGCTAGACACACTCCAGTCAAT GCCTGGAAAAGCGAAGAAATGCCCCTATGGGCTTCGCATATGACACCCCGCTGTTTTGACTCA CCCGAAGCCAGACACGCCATAAGGTCGCTCACAGAGCGGCTTTACATCGGGGGCCCCCTG ACTAATTCTAAAGGGCAGAACTGCGGCTATCGCCGGTGCCGCGAGCGGTGTACTCACG AAGCTCCAGGACTGCACGATGCTCGTATGCGGAGACGACCTTGTCGTTATCTGTGAAAGC GACGTCCGGAACCTATCCAGCAAGGCCGTTAACCACATCCGCTCCGTGTGGAAGGACTTG CTGGAAGACACTGAGACACCAATTGACACCACCATCATGGCAAAAAATGAGGTTTTCTGC 8 4 8 6 8 6 8 6 8 6 8 6 8 6 8 6

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GENERLSAGGRPVLFYKTDLSGALNELQDERARLSWLATTGVPCAAVLDVVTBA
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SDLYLVTRHADV1PVRRRGDSRGSLLSPRPVSYLKGSSGGPLLCPSGHAVG1FRAAVC
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YAAQGYKVLVLNPSVAATLGFGAYMSKAHGIDPNIRTGVRTITTGAPITYSTYGKFLA
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PNI EEVALSSTGEI PFYGKAI PIETI KGGRHI.I FCHSKKKCDEI.AAKI.SGI.GI.NAVAY
YRGLDVSVI.PTSGDVI VVATDALMTGFTGDFDSVI DCNTCVTQTVDFSI.DPTFTI ETT
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WARPDYNPPILLESWKDPDYVPPVVHGCPLPPAKAPPIPPRRKRTVVLESSTVSSALA
ELATKTRGSSESSAVDSGTATASPDQPSDDGDAGSDVESYSSMPPLEGEPGDPDLSDG
SWSTVSEEASEDVVCCSMSYTWTGALITPCAARETKLPINALSNSLLRHHNLVYATTS
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EASLRAFTEAMTRYSAPPGDPPKPEYDLELITSCSSNVSVAHDASGKRVYYLTRDPTT
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VRARLLSQGGRAATGGKYLFNWAVRTKLKLTPIPAASQLDLSSWFVAGYSGGDIYHSL
                                                                   'note="Neomycin-selectable bicistronic subgenomic
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/note="internal ribosome entry site (IRES)"
                                                                                                                                                                                                                                                                                                                                                                                                                                                 organism="Encephalomyocarditis virus"
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'product="core-neo fusion protein"
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join(1. .341,1801. .7758)
/organism="Hepatitis C virus"
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/db_xref="G1:5441835"
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'tissue_type="liver"
190. .1800
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/db_xref="taxon:12104"
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_xref="taxon:95363"
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AUZ42652.1 GI:5441834
COCE-NOS GENE; NS2 proteinase; NS3 gene; NS3
Proteinase/helicase; NS3/4A proteinase cofactor; NS4A gene; NS4B
GENE; NS4b protein; NS5a gene; NS5A phosphoprotein; NS5b gene; NS5B
GENE; NS4b protein; NS5a gene; NS5A protein; NS5B gene; NS5B
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Bartenschlager, R.
Direct Submission
Submitted (26-MAY-1999) Bartenschlager R., Institute for Virology, Johannes Gutenberg - University Mainz, Obere Zahlbacher Strasse 67, 55131 Mainz, GERMANY
Location/Qualifiers
                                                                                                                                                                                                                                                                                                                                                  TCATGCCTCAGGAAACTTGGGGTACCGCCCTTGCGAGTCTGGAGACATCGGGCCAGAAGT
                                                                          rcardccrcadgaaacrrdgggraccccrrdcgagrcrdgagacarcgggccagaagr
                                                                                                                                                                                                                        GrccGcGcTAGGCTACTGTCCCAGGGGGGGGCTGCCACTTGTGGCAAGTACCTCTTC
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/mol_type="other RNA"
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B & B & B ö g 6 6 6 6 6 ACCCGCGTGGAGTCAGAAAAAAAGTAGTAATTTTGGACTCTTTCGAGCCGCTCCAAGCG CCTCGAGCGATGCCCATATGGGCACGCCCGGATTACAACCCTCCACTGTTAGAGTCCTGG CTGTGCGGGTGGCTGCTGAGAAGTACGTGAGGGTTACGCGGGTGGGGGATTTCCACTAC TTCACAGAAGTGGATGGGGTGCGGTTGCACAGGTACGCTCCAGCGTGCAAACCCCTCCTA CGGGAGGAGGTCACATTCCTGGTCGGGCTCAATCAATACTGGTTGGGTCACAGGTCCCA ATTAACGCGTACACCACGGGCCCTGCACCCTCCCCGGCGCCAATTATTCTAGGGCG CGATTGCCGGGAGTCCCCTTCTTCTCATGAAGGGGGTACAAGGGAGGAGTCTGGCGGGGGGCCCTTCTTCTCATGTCAACGTGGGTACAAGGGAGTCTGGCGGGGGGCCCTTCTTCTCATGTCAACGTGGGTACAAGGGAGTCTGGCGGGGGC GACGCCATCATCCAAACCACCTGCCCATGTGGAGCACAGATCACCGGACATGTGAAAAAC GATTGGATATGCACGGTGTTGACTGATTTCAAGACCTGGCTCCAGTCCAAGCTCCTGCCG CAGTGGATCAACGAGGACTGCTCCACGCCATGCTCCGGGTCGTGGCTAAGAGATGTTTGG TTCGCTTCGCGGGGTAACCACGTCTCCCCCACGCACTATGTGCCTGAGAGCGACGCTGCA

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CCTCAGGCCGTGATGGGCTCTTCATACGGATTCCAATACTCTCCTGGACAGCGGGGTCGAGCTCTCAGGCCGGGTCGAGCCTCTTCATACGGATTCCAATACTCTCCTGGACAGCGGGTCGAG

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120 120 240 300 360 360 420 468 480 528 540 588 009 648 99 708 720 768 780 828 840

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949 ATGGCGATGCCTGCTTGCCGAATATCATGGTGGAAAATGGCCGCTTTTCTGGATTCATCG 1008
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                                                                                                                        Gaps
                                                                                                                        12;
                                                                        Length 8001;
                                                                                                                        Indels
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                                                                                 DB 12;
                                                                                                                          0; Mismatches
                                                                                 Score 7967;
Pred. No. 0;
                                                                                 99.78;
/gene="NS5B"
7771. .8001
                                                                                                                               Conservative
                                                                                                          Similarity
                                                                                                                                 Matches 7989;
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SWSTYGEEASEDVVCCSMSYTWTGALITPCAAEFTKLPINALSNSLLRHHNLVYATTS
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FGYGAKDVRNLSSKAVNHIRSYWKDLLEDTETPTDTTINAKNEVFCVQPEKGGRKPAR
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GACYSIEPLDILPQIIQRLHGLSAFSLHSYSPGEINRVASCLRKLGVPPLRVWRHRARS
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NGRFGGFTDCGRLGVADRYQDIALATRDIAEELGGEWADRFLVGTAAPDSQRIAFY
RLLDEFF"
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FAYDTRCFDSTVTENDIRVEESIYQCCDLAPEARQAIRSLTERLYIGGPLTNSKGQNC
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BASLRAFTEAMTRYSAPPGDPPKPEYDLELITSCSSNVSVAHDASGKRVYYLTRDPTT
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/note="internal ribosome entry site (IRES)"
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8709. .3870
gene="NS4A"
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5995. .7767
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3709. .3870
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3871 .4653
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Bartenschlager,R.
Hepatitis C virus culture system
Patent: US 6630343-A 16 07-OCT-2003;
Location/Qualifiers
                                                                                            Score 7947.8;
Pred. No. 0;
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/mol_type="genomic
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	TGTCGAGCTGCGGAAGCTCCACGTAATACCCTCACATGTTATTTGAAGGCCGCTGCGGCCTGCGAGCTGCGAGCTGCGAGCTGCAGATGTTATTTGAAGGCCGCTGCGGCCTGCGGCCTGCGAGCTGCAGAGCTCCAGGACTCCACGAGCCGAGCCACGAGCCGAGCCTTGTCGTTATTTGCGAGCACGACCTGCTTGTTGTTGTTGCGAGCCCCTTGTCGTTTGTCGAGCCGGGCGCGCCTTGTTGTTGTTGAGCGCGGGCGCCTTGTTGTTGTTGTTGTTGAGCGCGGGGCGCGCCTTCCGCCCCTTGTTGTTGTTGTTGTT	7081 ATACATCATCATCATCATCATCATCATCATCATCATCATC	7309 CTAGATTGTCAGATCTACGGGGCCTGTTACTCCATTGACCTTGACCTCCGGTC 7368

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3349 TACCTAAACACCAGGGTTGCCCGTCTGCCAGGACCATCTGGAGTTCTGGCAGAGCGTC 3408 3361 TACCTAAACACCACAGGGCTGCCCGTCTGCCAGGACCATCTGGAGTTCTGGCAGAGCGTC 3408 3409 TTACAGGCCTCACCCACTACACACCATTTCTTGTCCCAGACTTCTGGGAGGCGTC 3420 3409 TTACAGGCCTCACCCACATAGACGCCCATTTCTTGTCCCAGACTAGCAGGAGGCTC 3420 3411 TTACAGGCCTCACCCACATAGACGCCCATTTCTTGTCCCAGACTAGCAGGAGGC 3480 3421 TTACAGGCCTCACCCACATAGACGCCCATTTCTTGTCCCAGACTAGGCAGGAGAC 3480 3481 AACTTCCCCTACCTGGTAGCATACCAGGCTACGGTGTGCCCAGGGCTCAGGCTCCACCT 3528 3481 AACTTCCCCTACCTGGTAGCATACCAGGCTACGGCTCCAGGCTCCACCT 3540 3529 CCATCGTGGGCACAATGTGGAAGTCTCTCATACGCCTAGGCTACGCTCCACCT 3540 3541 CCATCGTGGGACCAAATGTGGAAGTCTCTCATACGCCTAACGCTACCACCCC 3588 3541 CCATCGTGGGAACCAAATGTGGAAGTCTCTCATACGCTTAAAACCTACACCACCA	CAAAACGAGTTACTACCACACACCCATA 36	ATTCCCGACAGGAAGTCCTT 38	400 402 406 408 412	424 424 426 426 426 426	
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6961 ATCTGTGAAAGCGCGGGACCCAAGAGGACGAGCCTACGGGCCTTCACGGAGGC 7009 ATGACTAGATACTCTGCCCCCCTGGGACCCGCCCAAACCAGAATACGACTTGAGTT [7141 TATCTCACCGTGACCCACCACCCCTTGCGGGCTGCGGGACACCAGCTAGACAC 7200 7189 ACTCCAGTCAATTCTGGCTAGGCAACATCATCATGTGGGCCCCCCCTTGTGGGCAAGG 7248	9 CTAGATTGTCAGAGATCTACGGGGCCTGTTACTCCATTGAGCCACTTGACCTACCT	7429 AATAGGGTGGCTTCATGCGAAACTTGGGGTACCGCCCTTGCGAGACAT 7488	7561 AAGTACCTCTTCAACTGGGCAGTAAGGACCTCAACTCACTC	7789 CCATCCTGTTTTTTCCTTTTTTTTTTTTTTTTTTTTTTT
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RESULT 13 AX036261

PAT 16-NOV-2000 Hepatitis C virus Hepatitis C virus Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae; Hepacivirus. 1; TCTTCACGCAGAAAGCGTCTAGCCATGGCGTTAGTATGAGTGTCGTGCAGGCCTCCAGGAC 120 360 121 CCCCCTCCCGGGAGGCCATAGTGGTCTCTGCGGAACCGGTGAGTACACCGGAATTGCCAG 180 240 240 241 GCGAGACTGCTAGCCGAGTAGTGTTGGGTCGCGAAAGGCCTTGTGGTACTGCCTGATAGG 300 408 409 ACGCAGATTCTCCGGCCGCTTGGGTGGAGGCTATTCGGCTATGACTGGGCACAACAGA 468 420 528 09 GCCAGCCCCCGATTGGGGGGGGACACTCCACATAGATCACTCCCCTGTGAGGAACTACTG 60 540 TIGICAAGACCGACCTGICCGGTGCCCTGAATGAACTGCAGGACGAGGCAGGGGGTAT 588 900 648 099 708 720 GCCAGCCCCGATTGGGGGGGACACTCCACCATAGATCACTCCCCTGTGAGGAACTACTG 181 GACGACCGGGTCCTTTCTTGGATCAACCCGCTCAATGCCTGGAGATTTGGGGGGTGCCCC 301 GIGCTIGCGAGTGCCCCGGGAGGTCTCGTAGACCGTGCACCATGAGCACGAATCCTAAAC 481 CAATCGGCTGCTCTGATGCCGCCGTGTTCCGGCTGTCAGCGCAGGGGCGCCCCGGTTCTTT 361 CTCAAAGAAAAACCAAACGTAACACCAACGGCGCGCCCATGATTGAACAAGATGCTTGC CTCAAAGAAAAACCAAA------GGGCGCGCCCATGATTGAACAAGATGGATTGC CAATCGGCTGCTCTGATGCCGCCGTGTTCCGGCTGTCAGGGGGGGCGCCCGGTTCTTT CGTGGCTGGCCACGACGGGCGTTCCTTGCGCAGGTGTGTGAACGTTGAAGCGG 12; Gaps DB 6; Length 8001; linear 16; Indels DNA Location/Qualifiers
1. 8001
/organism="Hepatitis C virus"
/db_xref="taxon:11103" Bartenschlager, R.D. Hepatitis c virus cell culture system Patent: EP 1043399-A 10 11-OCT-2000; BARTENSCHLAGER RALF DR (DE) ery Match 99.4%; Score 7941.4; st Local Similarity 99.7%; Pred. No. 0; tches 7973; Conservative 0; Mismatches AX036261 8001 bp Sequence 10 from Patent EP1043399. AX036261 GI:11225877 Н 19 181 361 469 649 529 589 601 199 US INITION ESSION SION WORDS RCE RGANISM source SRENCE URNAL Z 경 유

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Best Local Similarity 100.0%; Pred. No. 0;
Matches 7613; Conservative 0; Mismatches
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Reporter-selectable hepatitis c virus replicon
Patent: WO 03091439-A 2 06-NOV-2003;
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6558 7720 3	aticcaatacteteceggacaggegegetegagtecegegaatgeceggaaggeagga 6617 	음 ò	8800 TGCTGG1 7698 CATGTGC
6618	ATGCCCTATGGGCTTCGCATATGACACCGGCTGTTTTGACTCAACGGTCACTGAGAATGA 6677	2 d d	0 0
6678	CATCOGIGITGAGGAGTCAATCTACCAATGTTGTGGCTTTGGCCCCCGAAGCCACAGGC 6737	2 d 6	
	CATAAGGTCGCTCACAGAGCGGCTTTACATCGGGGGCCCCTGACTAATTCTAAAGGGCA 6797 	3 A S	
6798	GAACTGCGGCTATCGCCGGTGCCGCGAGCGGTGTACTGACGACCAGCTGCGGTAATAC 6857 	; 음 &	0 00
6858	CCTCACATGTTACTTGAAGGCCGCTGCGGCTGTCGAGCTGCGAAGCTCCAGGACTGCAC 6917 		
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7218	CATCATGTATGCGCCCACCTTGTGGGCAAGGATGCTGATGACCTGATTCTTCTCCCAT 7277 		
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7338	CTCCATTGAGCCACTTGACCTACGACATCATTCAACGACTCCATGGCCTTAGCGCATT 7397		
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7458	TGGGGTACCGCCCTTGCGAGTCTGGAGACATCGGGCCAGAAGTGTCCGCGCTAGGCTACT 7517		
7518	B GTCCCAGGGGGAGGGCTGCCACTTGTGGCAAGTACCTCTTCAACTGGGCAGTAAGGAC 7577 		

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search completed: November 2, 2004, 01:52:47
Job time : 21431 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.
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OM nucleic - nucleic search, using sw model

November 1, 2004, 17:15:25; Search time 398 Seconds (without alignments) 14272.926 Million cell updates/sec Run on:

US-10-005-469-1 7992

Perfect score:

1 gccagccccgattgggggc......tctctcgcagatcaagtact 7992 Sequence:

IDENTITY_NUC Gapop 10.0 , Gapext 1.0 Scoring table:

824507 seqs, 355394441 residues Searched:

1649014 Total number of hits satisfying chosen parameters:

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

/cgn2_6/ptcdata/1/ina/5A_COMB.seq:*/cgn2_6/ptcdata/1/ina/5B_COMB.seq:*/cgn2_6/ptcdata/1/ina/6A_COMB.seq:*/cgn2_6/ptcdata/1/ina/6B_COMB.seq:*/cgn2_6/ptcdata/1/ina/PCTUS_COMB.seq:*/cgn2_6/ptcdata/1/ina/PCTUS_COMB.seq:*/cgn2_6/ptcdata/1/ina/PCTUS_COMB.seq:*/ Issued_Patents NA:* Database :

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

STRAMMITES

	Description		9 6	, ,	acrembac	Semiented S	Seguence 29		Sequence 13, Appl	Semionos 24	Company 24,	Company of	Seguence 25,	1	٥١	'n.	4.	Sequence 1, 1	Sequence 25,	Sequence 19,	31,		31.	, [Cogramo 21		7	Sednence 1, Appli	ı,	-
SUMMARIES	ID	US-09-539-601-10	US-09-539-601-7	US-09-539-601-22	A1-109-539-60-SU	US-09-539-601-28	US-09-539-601-4	US-09-539-601-13	US-10-029-907-1	US-10-029-907-24	US-10-029-907-7	US-10-029-907-25	US-10-029-907-2	US-10-029-907-6	US-10-029-907-5	US-10-029-002	TIC-00-620 CT 1	T-109-238-60T-T	US-09-539-601-25	US-09-539-601-19	US-09-539-601-31	US-09-014-416-4	US-08-324-977-31	US-08-384-616-31	US-08-904-686A-3	US-09-315-850-31	TIC-000-224 077 1	176-08-354-977	0 0	US-08-904-686A-1
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5218	5218	5211.8	5175.2	5175.2	5175.2	5175.2	5175.2	5175.2	5175.2	5175.2	4103.8	4102.2	4097.8	4097.8	4094.2	4094.2	4021.4	
28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	

ALIGNMENTS

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FARTURE:

LOCATION: (1)...(341)

OTHER INFORMATION: construct I377/NS3-3'/wt

FRATURE:

NAME/KEY: CDS

LOCATION: (342).. (1181)

OTHER INFORMATION: hepatitis C virus core-neomycin phosphotransferase
FRATURE:

NAME/KEY: RBS

LOCATION: (1190)...(1800)

OTHER INFORMATION: internal ribosome entry site from
OTHER INFORMATION: encephalomyocarditis virus
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     AUTHORS: Lohman, Volker
AUTHORS: Krner, Frank
AUTHORS: Krner, Frank
AUTHORS: Krein, Jan-Oliver
AUTHORS: Herian, Ulrie
AUTHORS: Herilann, Lorenz
AUTHORS: Bartenschlager, Ralf
TITLE: Replication of subgenomic hepatitis c virus RNAs in a
TITLE: hepatoma cell line
VOUNDE: 285
                                                                                     APPLICANT: Bartenschlager, Ralf FW
TITLE OF INVENTION: Hepatitis C Virus Cell Culture System
FILE REFERENCE: all sequences
CURRENT APPLICATION NUMBER: US/09/539,601C
GURRENT FILING DATE: 2001-08-30
EARLIER PILING DATE: 1999-04-03
NUMBER OF SEQ ID NOS: 51
SOFTWARE: Patentin Ver. 2.1
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             FEATURE:
NAME/KEY: CDS
LOCATION: (1801)..(7758)
OTHER INFORMATION: hepatitis C virus N83 - 5B
                      Sequence 10, Application US/09539601C Patent No. 6630343
                                                                                                                                                                                                                                                                                                                                                      TYPE: DNA ORGANISM: Hepatitis C virus
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PUBLICATION INFORMATION:
                                                                  GENERAL INFORMATION:
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NAME/KEY: 3'UTR
LOCATION: (7759).
US-09-539-601-10
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LENGTH: 7989
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Ouery Match Query Match Query Match Query Match Dest Local Similarity 100.0%; Score 7989; DB 4; Length 7989; Best Local Similarity 100.0%; Pred. No. 0; Matches 7989; Conservative 0; Mismatches 0; Indels 0; Gaps 0; Query Matches 7989; Conservative 0; Mismatches 0; Indels 0; Db		
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	7921 7921 7981 7981 7981 7981 7981 7981 7981 798	ORGANISM: Hepatitis C virus FRATURE: NAME/KEY: NAME/KEY: COATION: (1)(341) COTHER INFORMATION: construct 1389/NS3-3'/wt FEATURE: NAME/KEY: COATION: (342)(1193) COTHER INFORMATION: hepatitis C virus core-neomycin FEATURE: FEATURE: COATION: (1202)(1812) COTHER INFORMATION: internal ribosome entry site from FEATURE: NAME/KEY: CDS COCHER INFORMATION: encephalomyocarditis virus FEATURE: NAME/KEY: CDS COCHER INFORMATION: hepatitis C virus nonstructural proteins FEATURE: NAME/KEY: ADS LOCATION: (1813)(7770) COTHER INFORMATION: hepatitis C virus nonstructural proteins NS3-5B FEATURE: NAME/KEY: 3'UTR LOCATION: (7771)(8001)
6421 GTCCAACCAGAGAAGGGGGCCGCAAGCCAGCTTATCGTATTCCTATTTGGGG 6480 6481 GTTCGTGTGTGCGAGAAATGGCCTTTACGATGGTCTCCACCCTCCAGGCGTG 6540 6481 GTTCGTGTGTGCGAGAAATGGCCTTTACGATGGTCTCCACCCTCCTCAGGCGGTG 6540 6481 GTTCGTGTGTGCGAGAAATGGCCTTTACGATGTGTTCCACCCTCCTCAGGCGGTG 6540 6541 ATGGGCTCTTCATACGGATTCCAATACTCTCCTGGACAGCGGTCGAGTTCCTGGTGAAT 6600 6541 ATGGGCTCTTCATACGGATTCCAATACTCTCTGGACAGCGGTCGAGTTCTTGGTGAAT 6600 6601 GCCTGGAAAGGAAAAAGGCTTCCAATACACTCGGACAGGGGTCCAGGTTTTTGACTCA 660 6601 GCCTGGAAAGGAAAAGGCTTTGAGGTTTCGAAATGACACCCGCTGTTTTTGACTCA 660 6601 GCCTGGAAAGGAAAAGACCCTATGAGGCTTCGCATATGACACCCGCTGTTTTTGACTCA 660 6601 GCCTGGAAAGGAAAGACATCCGTGTTGAAATGCCAAATGAAACCGCTTTTGACTCA 660 6601 GCCTGGAAAGGAAAGACATCCGTGTTTGAAATGACACCCGCTGTTTTTGACTCA 660 6601 GCCTGGAAAGGAAAGACATCCGTGTTTGAAATTTACATCGGCGGGCCCCCTG 6780 6721 CCCGAAGCCAGACAGGCCATAAGGTCAAACTCAAGGGGGGCCCCCTG 6780 6722 CCCGAAGCCAGACAGGCCATAAGGTCAAACGGGGGCCCCCTG 6780 6724 ACTAATTCTAAAGGGCATACGCGCTATCGCGGGGCCCCCTG 6780 6781 ACTAATTCTAAAGGGCAAACTGCGGCTATCGCGGGGCCCCCTGGGGGCCCCCTG 6780 6781 ACTAATTCTAAAGGGCAAACTGCGGCTATCGCCGGGCCCCCCGGGGGCCCCCTGGGGGGCCCCTGGGGGG		7201 †CÉTGGCTAGGCAACATCATGATGGGCCCACCTTGTGGGCAAGGATGTCCTGTG 7261 ACTCATTTCTTCTCCTCTTCTAGCTCGGAACAACATGAAAAGCCCTAGATTGTCAG 7261 ACTCATTTCTTCTCCTCTTTAGCTCAGGAACAACTTGAAAAAGCCCTAGATTGTCAG 7321 ATCTAGGGGGCCTGTTACTCCATTGACACTTGAAAAAGCCCTAGATTGTCAG 7321 ATCTACGGGGCCTGTTACTCCATTGACACTTGACACTCAGATCATTCAACGACTC 7321 ATCTACGGGGCCTGTTACTCCATTGACACTTGACTACTCAGATCATTCAACACACTC 7381 CATGGCCTTAGCGCATTTTCACTCCATTGACTCTCCCAGATCATTCAACACACTC 7381 CATGGCCTTAGCGCATTTTCACTCCATTGACTTCTCCCAGATCAATAGGGTGCCT 740 TATGTCACGGGAACTTGAGCTACTCCTCCAGATGAGACTCAGGTGCCT 741 TCATGCCTCAGGAAACTTGGGGTACCGCCCTTGCGAGTCTGCGAGACATCGGGCCAGAAGT 7501 GTCCGCGCTAGGGTACCGCCCTTGCGGTACCTCTCGAGACATCGGGCCAGAAGT 7501 GTCCGCGCTAGGCTACCGCCCTTGCGGGGCCACAAGTACCTCTTC 7501 GTCCGCGCTAGGCTACTGCCCAGGGGGGGGGCCCACTTGTCGCCAAGTACCTCTTC 7501 GTCCGCGCTAGGCTACTGCCCAGGGGGGGGGCGCCACTTGTCGCCAAGTACCTCTTC 7501 GTCCGCGCTAGGCTACTGCCCAGGGGGGGGCGCCCATTGTCGCCAAGTACCTCTTC 7501 GTCCGCCCTAGGCTACTGCCCAGGGGGGGGGCAGGACCATTGTCGCCAAGTACCTCTTC 7501 GTCCGCCCTAGGCTACTGCCCAGGGGGGGCAGGACCACTTGTCGCCAAGTACCTCTTC 7501 GTCCGCCCTAGGCGTACCTGCCCAGGGGGGCAGCACTTGTCGCCAAGTACCTCTTC 7501 GTCCGCCCTAGGGGGAGGGGCCCCATTGTCGCCAAGTACCTCTTC 7501 GTCCGCCCTAGGGGGGGGGGGGCCCACTTGTCGCCAAGTACCTCTTC 7501 GTCCGCCTAGGGGTACCTGCCCAGGGGGGCAGCAAGTACCTCTTC 7501 GTCCGCCTAGGGCTACCTGCCCAGGGGGCACCAAGTACCTCTTC 7501 GTCCGCCTAGGCTACCTGCCCCTTGCCCCATTGTCGCCAAGTACCTCTTC 7501 GTCCGCCTAGGCGTACCTGCCCCAGGGGGCAGCAGTACCTCTTC 7501 GTCCGCCTAGGGGGAGGGGCACAGTACCTCTTC 7501 GTCCGCCTAGGGGGAGGGCACACTTGTCGCCCAAGTACCTCTTC 7501 GTCCGCCTAGGGGGAGGGGCACCCAGGGGGGCACAGTACCTCTTC 7501 GTCCGCCTAGGCGTACCTTCTCCCCAGGGGGGCACTAGCTCTCTCCTCTCTCT

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    PUBLICATION INFORMATION:

AUTHORS: Lohmann, Volker

AUTHORS: Krner, Prank

AUTHORS: Herian, Ulrike

AUTHORS: Theilmann, Lorenz

AUTHORS: Theilmann, Lorenz

AUTHORS: Partenschlager, Ralf

ITILE: Replication of subgenomic hepatitis c v

TITLE: Replication of subgenomic hepatitis c v

TITLE: Replication of subgenomic hepatitis c v

TITLE: Replication

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Best Local Similarity 99.9
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NAME/KEY: RBS

LOCATION: (1202)..(1812)

GTHER INFORMATION: internal ribosome entry site from OTHER INFORMATION: encephalomyocarditis virus
FEATURE:
NAME/KEY: CDS

LOCATION: (1813)..(7770)

GTHER INFORMATION: hepatitis C virus nonstructural proformation of cell culture-adapted clone no. 5
FEATURE:
NAME/KEY: 5'UTR

LOCATION: (7771)..(8001)
US-09-539-601-22
                                                   no.
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                                                                                        Score 7949.4;
Pred. No. 0;
0; Mismatches
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Best Local Similarity 99.7%;
Matches 7978; Conservative
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Db 7381 ATTCAACGACTCCATGGCGAATTTTCATCCTCCAAGTTACTCCCAGGGGGGGG	RESULT 4 US-09-59-601-16 Sequence 16, Application US/09539601C TITLE OF INVENTION: Hepatitis C Virus Cell Culture System FILE REPRENCE: all sequences CURRENT RELING DATE: 2000-08-30 BARLIER APPLICATION UNMERE: 199 15 178.4 GERMANY MUMBER OF SEQ ID NOS: 51 SOFTWARE: Patentin Ver. 2.1 SOFTWARE: Patentin Ver. 2.1 SOFTWARE: Patentin Ver. 2.1 SOFTWARE: Patentin C Virus NAME/KEY: S'UTR COCATION: (1)(341) COCATION: (1)(342) COCATION: (1)(342) COCATION: (1342) COCATION: (1342)(1193) COTHER INFORMATION: hepatitis C virus core-neomycin phosphotransferase COTHER INFORMATION: fusion protein FRATURE: NAME/KEY: RBS

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                                                                                                                       Gaps
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                                                     NS3
         LOCATION: (1202)...(1812)

OTHER INFORMATION: internal ribosome entry site from OTHER INFORMATION: internal ribosome entry site from OTHER INFORMATION: encephalomyocarditis virus
FEATURE:
NAME/KEY: CDS
LOCATION: (1813)...(7770)
OTHER INFORMATION: of cell culture adapted clone no. 19
FEATURE:
NAME/KEY: 3'UTR
NOME/KEY: 3'UTR
LOCATION: (1771)...(8001)
US-09-539-601-28
                                                                                                         Length
                                                                                                                       Indels
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                                                                                                          Score 7941.4;
Pred. No. 0;
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Best Local Similarity 99.7%;
Matches 7973; Conservative 0
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Db 7381 ATTCAACGACTCCATGGCGCATTTTCACTCCATAGTTACTCTCCAGGTGAGATC 7440	QY 7489 CGGGCCAGAAGTGTCCCGCGCTAGGCTACTGTCCCAGGGGGGGG	Oy 7549 AAGTACCTCTTCAACTGGGCAGTAAGGACCAAGCTCAAACTCCCAATCCCGGCTGCG 7608	Qy 7609 TCCCAGTTGGATTATCCAGCTGGTTGGTTGCTACAGCGGGGGAGCACATATATCAC 7668	Oy 7669 AGCCTGTCTGTGCCGACCCGCTGGTTCATGTGGTGCTACTCCTACTTTTGTAGGG 7728	Qy 7729 GTAGGCATCTACTCCCCAACOGATGAACGGGGAGCTAAACACTCCAGGCCAATAGG 7788	Oy 7789 CCATCCTGTTTTTTTCTTTTTTTTTTTTTTTTTTTTTTT	QY 7849 TITCLCCTTTTTTTTTTTCCTTTTTCCTTTGGGGGCTCCATCTTAGC 7908 DD 7861 TITCTCCTTTTTTTTTTTTTTTTTTTTTTTCCTTTTTTTT	OY 7909 CCTAGTCACGGCTAGCTGAAAGGTCCGTGAGCCGCTTGACTCCAGAGAGTGCTGATAC 7968	OY 7969 TGGCCTCTCTGCAGATCAAGT 7989	RESULT 6 US-09-539-601-4 ; Sequence 4, Application US/09539601C	; Patent No. 6630343 ; GENERAL INFORMATION: ; APPLICANT: Bartenschlager, Ralf FW ; TITLE OF INVENTION: Hepatitis C Virus Cell Culture System	NAMA MA		; LENGTH: 8637 ; TYPE: DNA ; ORGANISM: Hepatitis C virus ; FEATURE:	; NAME/KEY: 5'UTR ; LOCATION: (1)(341) ; CTHER INFORMATION: construct 1377/NS2-3'/wt ; FEATURE:	: NAME/KEY: CDS : LOCATION: (342)(1181) : COTHER INFORMATION: HCV core-neomycin phosphotransferase fusion : OTHER INFORMATION: protein	; FEATURE: ; NAME/KEY: RBS ; LOCATION: (1190)(1800)
	CGTATTC 	6469 CCAGATTIGGGGGTTCGTGTGTGCGAAAAATGGCCCTTTACGATGTGTCTCCACCTC 6528 	6529 CCTCAGGCCGTGATGGGCTCTTCATACGGATTCCAATACTCTCCTGGACAGCGGGTCGAG 6588 	6589 TTCCTGGTCAATGCCTGGAAAGCGAAGAATGCCCTATGGGCTTCGCATATGACACCCGC 6648 	6649 TGTTTTGACTCAACGGTCACTGAGAATGACATCCGTGTTGAGGAGTCAATCTACCAATGT 6708 	6709 TGTGACTTGGCCCCCGAAGCCAGACAGGCCATAAGGTCGCTCACAGAGCGGCTTTACATC 6768 	6769 GGGGCCCCCTGACTAATTCTAAAGGGCAGAACTGCGGCTATCGCCGGTGCCGGGCGGG	6829 GGTGTACTGACGACCAGCTGCGGTAATACCCTCACATGTTACTTGAAGGCCGCTGCGGCC 6888	6889 TGTCGAGCTGCGAAGCTCCAGGACTGCACGATGCTCGTATGCGGAGACGACCTTGTCGTT 6948 	6949 ATCTGTGAAAGCGCGGGGACCCAAGAGGACGAGCCGTACGGGCCTTCACGGAGGCT 7008 	7009 ATGACTAGATACTCTGCCCCCCTGGGGACCCGCCCAAACCAGAATACGACTTGGAGTTG 7068 	7069 ATAACATCATGCTCCTCCAATGTCAGTCGCGCACGATGCATCTGGCAAAAGGGTGTAC 7128 	7129 TATCTCACCGGGGCCCCCCCCCCCCTGGGGGGCTGCGTGGGAAGACAGCTAGACAC 7188 	7189 ACTCCAGTCAATTCCTGGCTAGGCAACATCATGTATGCGCCCACCTTGTGGGCAAGG 7248 	7249 ATGATCCTGATGACTCATTTCTTCCATCCTTCTAGCTCAGGACAACTTGAAAAAGCC 7308 	7309 CTAGATIGICAGAICTACGGGGCCTGTTACTCCATIGAGCCACTTGACCTACCTGAGTC 7368 	7369 ATTCAACGACTCCATGGCCTTAGCGCATTTCACTCCATAGTTACTCTCCAGGGGGGAGATC 7428

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OTHER INFORMATION: internal ribosome entry site from OTHER INFORMATION: encephalomyocarditis virus FEATURE:
NAME/KEY: CDS
LOCATION: (1801)..(8406)
OTHER INFORMATION: hepatitis C virus NS2 - 5B
FEATURE:
NAME/KEY: 3. UPO
LOCATION: (8407)..(8637)
FUBLICATION INFORMATION:
AUTHORS: Lohamann, Volker
AUTHORS: Koch, Jan-Oliver
AUTHORS: Merian, Ulriker
AUTHORS: Heilmann, Lorenz
AUTHORS: Bartenschlager, Ralf
TITLE: hepatoma cell line
JOURNAL: Science
JOURNAL: Science
JOURNAL: 285
DESEC: 1.0.1
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Best Local Similarity 92.5
Matches 7989; Conservative
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                                                                                                                                    CCCCCCTCCCGGGAGAGCCATAGTGGTCTGCGGAACCGGTGAGTACACCGGAATTGCCAG
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                                                                   Gaps
                                                                 Indels 660;
                           Length 8649;
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0
                               DB 4;
                                                                       0; Mismatches
                           Score 7309;
Pred. No. 0;
                                   91.5%;
                                                                         Conservative
                                                      Similarity
                                 Query Match
Best Local Simi
Matches 7989;
US-09-539-601-13
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OTHER INFORMATION: construct 1389/NS2-3'/wt
FEATURE:
NAME/KEY: CDS
LOCATION: (342)...(1193)
OTHER INFORMATION: hepatitis c virus core-neomycin phosphotransferase
OTHER INFORMATION: fusion protein
                                                                                    CGATGAACGGGGAGCTAAACACTCCAGGCCAATAGGCCATCCTGTTTTTTTCCCTTTTTT
                                                                                                               LOCATION: (1813)..(8418)
LOCATION: (1813)..(8418)
COTHER INFORMATION: hepatitis C virus NS2 - 5B
FEATURE:
NAME/KEY: 3'UTR
LOCATION: (8419)..(8649)
PUBLICATION: NIPORMATION:
AUTHORS: Lohmann, Volker
AUTHORS: Kroer, Prank
AUTHORS: Kroer, Jan-Oliver
AUTHORS: Rech, Jan-Oliver
AUTHORS: Theilmann, Lorenz
AUTHORS: Perinschlager, Ralf
ITILE: Replication of subgenomic hepatitis c virus RNAs in a
TITLE: hepatoma cell line
JOURNAL: Science
VOLUME: 285
PAGES: 110-113
DATE: 1999-07-02
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; Patent No. 6630343
; GENERAL INFORMATION:
APPLICANT: Bartenschlager, Ralf FW
APPLICANT: Bartenschlager, Ralf FW
TITLE OF INVENTION: Hepatitis C Virus Cell Culture;
FILE REFERENCE: all sequences
CURRENT APPLICATION NUMBER: US/09/539,601C
CURRENT FILING DATE: 2001-08-30
EARLIER FILING DATE: 1999-04-03
; NUMBER FILING DATE: 1999-04-03
; NUMBER FILING DATE: 1990-04-03
; SOFTWARE: PatentIn Ver. 2.1
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NAME/KEY: RBS
LOCATION: (1202)..(1812)
COTHER INFORMATION: internal ribosome entry site
OTHER INFORMATION: encephalomyocarditis virus
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ORGANISM: Hepatitis C virus
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NAME/KEY: 5/UTR
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LENGTH: 8649
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US-09-539-601-13
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420 468

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1804 1803 2101 ATAACCAAAGTGCCGTACTTCGTGCGCGCACACGGGCTCATTCGTGCATGCTGGTG 2160 1804	2161 CGGAAGGTTGCTGGGGGTCATTATGTCCAAATGGCTCTCATGAAGTTGGCCGCACTGACA 2220 1804		1804 1803 2341 TGGGGGCAGACACCGCGGCGTGTGGGGACATCATCTTGGGCCTGCCCGTCTCCGCCCGC	1804GCGCCTATTACGGCCTACCCACAGCCTGCGCTGGAGGGCGGGGGGGCGCGACTC 2460 1804GCGCCTATTACGGCCTACTCCCAAAAGACGCGAGGCCTACTTGGCTGCATCATCACT 1860 2461 CTCGCGCCTATTACGGCCTACTCCCAACAGACGCCAAGCCTACTTGGCTGCATCATCACT 5520		1921 ACACAATCTTTCCTGGCGACCTGCGTCAATGGCGTGTGTTGGACTGTCTATCATGGTGCC 1980 	1981 GGCTCAAAGACCCTTGCCGGCCCAAAGGCCCCAATGACCCAAATGTACACCAATGTGGAC 2040 	2041 CAGACCTCGTCGGCTGGCAAGCGCCCCCCGGGGGGTTCCTTGACACCATGCACCTGC 2100	2101 GGCAGCTCGGACCTTTACTTGGTCACGAGGCATGCCGATGTCATTCCGGTGCGCGGGCGG				2341 CGGTCCCGGTCTTCACGACAACTCGTCCCCTCCGGCCGTACCGCAGACATTCCAGGTG 2400 3001 CGGTCCCCGGTCTTCACGGACAACTCGTCCCCTCCGGCCGTACCGCAGACATTCCAGGTG 3060		2401 GCCCAGGGTATAAGGTGCTTGTCCTGAACCCGTCGGTCGCGCCGCCTTGGGTTTCGGG 2520 3121 GCCCAAGGGTATAAGGTGCTTGTCCTGAACCGTCGTCGCCGCCACCCTAGGTTTCGGG 3180 2521 GCGTATATGTCTAAGGTGTTTCGGCCTCGTCGTCGCGCGCACCCTAGGTTTCGGG 3180
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	941 TCTACCGTAAGCGAGGCTAGTGAGGACGTCGTCTGCTCGATGTCCTACACATGG 600 	01 ACAGGGCCCTGATCACGCCATGCGCTGCGGAGGAAACCA 	21 AGC	21 CTGCGGCAGAA 	6181 GTGCTCAAGGAGTGAAGGCGAAGGCGTCCACAGTTAAGGCTAAACTTCTATCCGTGGAG 6240 		01	5=5	6421 GTCCAACCAGAGAAGGGGGGCCGCAAGCCAGCTCGCCTTATCGTATTCCCAGATTTGGGG 6480 	GTTCGTGTGT GTTCGTGTGT	541	601 GCCTG 261 GCCTG		81 0	81 41 41	6841 ACCAGCTGCGGTAATACCCTCACATGTTACTTGAAGGCCGCTGCGGCCTGTCGAGCTGCG 6900	
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CGACGGGCGTTCCTTGCGCAGCTGTGCTCGACGTTGTCACTGAAGCGGGAAGGGACTGGC 660
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                                                             GENERAL INC. 0/V00/13.

GENERAL INFORMATION:
GENERAL INFORMATION:
APPLICANT: BOEHERINGER INGELHEIM (CANADA) LTD.
TITLE OF INVENTION: SELF REPLICATING RNA MOLECULE FROM
TITLE OF INVENTION: HEPATITIS C VIRUS
FILE REFERENCE: 13/083
CURRENT APPLICATION NUMBER: US/10/029,907
CURRENT FILING DATE: 2001-12-21
PRIOR FILING DATE: 2000-12-22
NUMBER OF SEQ ID NOS: 25
SEQ ID NO 1
SEQ ID NO 1
LENGTH: 8639
TYPE: DNA 1
                                                                                                                                                                                                                                                                   DB 4;
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91.0%; Score 7273.8;
Best Local Similarity 92.2%; Pred. No. 0;
Matches 7976; Conservative 0; Mismatches
                                             ; Sequence 1, Application US/10029907; Patent No. 6706874
                                                                                                                                                                                                                               NAME/KEY: CDS
LOCATION: (1803)...(8408)
8641 AGATCAAGT 8649
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; NAME/KEY: CDS ; LOCATION: (1802)(8407) US-10-029-907-7	Query Match 90.7%; Score 7249.8; DB 4; Length 8638; Best Local Similarity 92.0%; Pred. No. 0; Matches 7961; Conservative 0; Mismatches 17; Indels 671; Gaps 3:	TICCCCTGTGAGGAACTACTG 60		CCAG 18 CCAG 18	OY 181 GACGACCGGGTCCTTTCTGGATCAACCGCTCAATGCCTGGAGATTTGGGCGTGCCCC 240	241 GCGAGACTGCTAGCCGAGTAGTGTTGGGTCGCGAAAGGCCTTGTGGTACTGCCTGATAGG	301	OY 361 CTCAAAGAAACCAAAGGGGCGCCATGATTGAACAAGATGGATTGCAGGCAG		r, r	541 ACCIGTCCGGTGCCCTGAATGAACTGCAGGACGAGGCAGCGGGGCTATCGTGGCTGGC	601	661		781 CATTCGACCACCAAGCGAAACATCGAGCGAGCAGGTACTCGGATGGAAGCCGGTC 	OY 841 TTGTCGATCAGGATGATGAGGAGGAGGGGCCTCGCGCCAGCCGAACTGTTCG 900	6	TGGTGGAAAATGGCCGCTTTTCTGGATTCATCGACTGTGGCCC

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90.7%; Score 7248.8; DB 4; Length 8638;
Best Local Similarity 92.0%; Pred. No. 0;
Matches 7960; Conservative 0; Mismatches 17; Indels 671;
                                                                                           GENERAL INFORMATION:
APPLICANT: BOEHRINGER INGELHEIM (CANADA) LTD.
TITLE OF INVENTION: SELF REPLICATING RNA MOLECULE FROM
TITLE OF INVENTION: HEPATITIS C VIRUS
FILE REFERENCE: 13/083
CURRENT FILING DATE: 201-12-21
PRIOR PLICATION NUMBER: 60/257,857
PRIOR PLICATION NUMBER: 60/257,857
PRIOR PLICATION NUMBER: 60/257,857
PRIOR PLICAND APPLICATION NUMBER: 60/257,857
SOFTWARE: FastSEQ for Windows Version 4.0
SOFTWARE: BOSE ID NOS: 25
LENGTH 8638
TYPE: DNA
TYPE: DNA
                                                                           Sequence 25, Application US/10029907
Patent No. 6706874
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; LOCATION: (1802)...(8407)
US-10-029-907-25
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Fatent No. 6706874
GENERAL INFORMATION:
APPLICANT BOENRINGER INGELHEIM (CANADA) LTD.
TITLE OF INVENTION: BELF REPLICATING RNA MOLECULE FROM
TITLE OF INVENTION: HEPATITIS C VIRUS
FILE REFERENCE: 13/083
CURRENT APPLICATION NUMBER: US/10/029,907
CURRENT APPLICATION NUMBER: 60/257,857
PRIOR FILING DATE: 2000-12-22
NUMBER OF SEQ ID NOS: 25
NUMBER OF SEQ ID NOS: 25
SOFTWARE: FastSEQ for Windows Version 4.0
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90.7%; Score 7246.8;
Best Local Similarity 92.1%; Pred. No. 0;
Matches 7966; Conservative 2; Mismatches
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NAME/KEY: CDS
LOCATION: (1802)...(8407)
NAME/KEY: variation
LOCATION: 6266
OTHER INFORMATION: r = a or g
NAME/KEY: variation
LOCATION: 4446
COTHER INFORMATION: r = a or g
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ORGANISM: HCV
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PRIOR FILING DATE: 2000-12-22
NUMBER OF SEQ ID NOS: 25
SOFTWARE: FastSEQ for Windows Version SEQ ID NO 6
LENGTH: 8638
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Best Local Similarity 92.0%;
Matches 7958; Conservative C
                                                                                 ... (8407)
                                                                      NAME/KEY: CDS
LOCATION: (1802)
                                               TYPE: DNA
ORGANISM: HCV
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Patent No. 6706874
GRNERAL INPORMATION:
APPLICANT: BOEHRINGER INGELHEIM (CANADA) LTD.
TITLE OF INVENTION: BEPRINGER REPLICATING RNA MOLECULE FROM
TITLE OF INVENTION: HEPATITIS C VIRUS
FILE REFERENCE: 13/083
CURRENT APPLICATION NUMBER: US/10/029,907
CURRENT FILING DATE: 2001-12-21
PRIOR APPLICATION NUMBER: 60/257,857
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                    19; Indels 671;
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Score 7245.6;
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TITTCCTCTTTTTTCCTT TTTCCTCTTTTTTCCTT TTTCCTCTTTTTT	Query Match 90.5%; Score 7229.8; DB 4; Length 8648; Best Local Similarity 91.9%; Part No. 0; Matches 7961; Conservative 0; Mismatches 17; Indels 681; Gaps 4; QV GCGCCCCCGATTGGGGGGGGGGGGGGGGGGGGGGGGGGG
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IGTACACCAATGTGGAC 2040 GCCCGTCTCCGCCCGC 2389 GCAGGGTGGCGACTC 2449 CCACGCGGCCTACGA 2269 GACCAAGGTTATCACC 2329 recardcardcrearg 2149 GITGGCCGCACTGACA 2209 SATCCCCCCCTCAAC 1969 CCACCCAGAGCTAATC 2029 ------ 1803 SGIGCICCAGGCIGGI 2089 GCGGTTTTCGTAGGT 1849 AGGCTCATATGGTGG 1909 ----- 1803 TTTTCCTTTGAAAA 1788 |||||||||||||||| |TTTCCTTTGAAAA 1789 ----- 1803

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SUMMARIES	CK284786 CK281519 CK287297 CK2877930 CK286377 CK288361 CK288361 CK288311 AQ38811 AQ38811 AQ44775 AQ44162 AQ398160 AQ44162 AQ398160 AQ44162 AQ398130 AQ398130 AQ397768 AQ397768	
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library, normalized, full-length...

/note="Wector: pCWNSport6.1; Site 1: EcoRI; Site 2: NotI;

/note="Wector: pCWNSport6.1; Site 1: EcoRI; Site 2: NotI;

supplier: RNA was isolated from Nicotiana benthamiana
tissues that include callus, roots from liquid culture
grown plants, heat-stressed leaves (38 C, 3 hr and 6 hr),

cold-stressed leaves (5 C 3 hr, 6hr), and pathogen
challenged leaves (Rseudomonas syringae pv tomato 12 hr;

Xanthomonas campestris pv campestris 12 hr, 18hr;

Pseudomonas syringae pv phaseolicola 18hr, and Xanthomonas
campestris pv vesicatoria 18hr). RNA was isolated from
these tissues and pooled in approximately equal molar
                                        Butaryota, Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Bukaryota, Viridiplantae; Streptophyta; eudicotyledons; core eudicots; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; asterids; lamiids; Solanales; Solanaceae; Nicotiana.

1 (baees 1 to 804)
Buell, C.R., Hart, A., Zismann, V., Karamycheva, S.A., Day, B., Graskawicz, B., Jin, H. and Baker, B.
Generation of EST sequences from Nicotiana benthamiana
                                                                                                                                                                                                                                                                                                                                                                                                                                                        /tissue type="abiotic and biotic stress-treated leaves, tissue type="abiotic and root tissue" | /lab host="DH10B-TonA" | /clone_lib="Nicotiana benthamiana mixed tissue cDNA | /clone_lib="Nicotiana benthamiana mixed tissue
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Contact: Robin Buell
The Institute for Genomic Research
5712 Medical Center Dr. Rockville, MD 20850, USA
Email: potato-arrayeligr.org
Clones can be requested from the University of Arizona Genomics
Institute via http://genome.arizona.edu/orders/ .
Seq primer: ATT TAG GTG ACA CTA TAG.
Location/Qualifiers
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                                      Nicotiana benthamiana
                Nicotiana benthamiana
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BST754233 Nicotiana benthamiana mixed tissue cDNA library, normalized, full-length Nicotiana benthamiana cDNA clone NBMC276 5'
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                                                                                 10.0%; Score 795.4; DB 7;
99.9%; Pred. No. 1.8e-174;
tive 0; Mismatches 1;
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challenged leaves (Pseudomonas syringae pv tomato 12 hr; Xanthomonas campestris pv campestris 12 hr, 18hr; Pseudomonas syringae pv phaseolicola 18hr; and Xanthomonas campestris pv vesicatoria 18hr). RNA was isolated from these tissues and pooled in approximately equal molar amounts."	Query Match 9.9%; Score 794; DB 7; Length 856; Best Local Similarity 100.0%; Pred. No. 3.8e-174; Matches 794; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	Qy         389         GATTGAACAAGATTGCACGCAGGTTCTCCGGCCGCTTGGGAGGCTATTCGG         448           Db         14         GATTGAAGAAGATGGATTGCACGCAGGTTCTCCCGGCCGCTTGGGTGGAGGAGGCTATTCGG         73	Oy 449 CTATGACTGGGCAGAACAATGGGCTGCTCTGATGCGGGGTGTTCGGGCTGTCAGC 508  14	Oy 509 GCAGGGGCCCGGTTCTTTTGTCAAGACCGACCTGCCGGTGCCCTGAATGAA	Qy         569         GGACGAGCGCGGCTATCGTGGCTACGGCCACGACGGCGTTCCTTGCGCAGCTGTGCT         628           Db         194         GGACGAGCGGCGTATCGTGGCTGGCCACGACGGCGTTCCTTGCGCAGCTGTGCT         253	Ay 629 CGACGITGICACTGAAGCGGAAGGGACTGGCTGCTATTGGGCGAAGTGCCGGGCAAGA 688							CGTGCTTTACGGTATGCCGGTCCCGATTGCAGCGCATCGCCTTCTATCGCTTCTGA 11  CGTGCTTTACGGTATGCCGGTCCCGATTGCAGCGCATCGCCTTCTATCGCTTCTTGA 11  CGTGCTTTAACGGTATCGCCGCTCCCCATTGCATCGCTAGCTA		RESULT 4 CK287930 LOCUS CK287930 DEFINITION EST750652 Nicotiana benthamiana mixed tissue cDNA library,
Qy         809 CGAGCGAGCACGTACTCGGATGGAAGCCGGTCTTGTCGATCAGGATGATCTGGACGAAGA 868           Db         422 CGAGCGACGTACTCGGATGGAAGCCGGTCTTGTCGATCAGGATCTGGACGAAGA 481           Qy         869 GCATCAGGACGTCGCCCAGCCCAGCCCAGCCCCAGGCTCCGACGC 928           Db         482 GCATCAGGGGCTCGCCCAGCCCAAGCCCGAGCTTGTCTAAGCCCCATGCCCAACGCTGACGC 641						RESULT 3 CK287297	LOCUS  CX287297  DEFINITION EST750019 Nicotiana benthamiana mixed tissue cDNA library, normalized, full-length Nicotiana benthamiana cDNA clone NBMB815 5' end, mRNA sequence.	ACCESSION CK287297 TVERSION CK287297.1 GI:39863696 KEYWORDS EST. SOURCE Nicotiana benthamiana	5	REFERENCE 1 (bases 1 to 856) AUTHORS Buell, C.R., Hart, A., Zismann, V., Karamycheva, S.A., Day, B., Staskawicz, B., Jin, H. and Baker, B. TITLE Generation of EST seminores from	9712 Medical Center Dr. Rockville, MD 20850, USA Email: potato-array@tigr.org clones can be requested from the University of Arizona Genomics Institute via httn://denome ariana ani/waxa/	Seq primer: ATT TAG GTG ACA CTA TAG.  FEATURES  Location/Qualifiers  1. 856	/mol_type="mknh" /db_xref="taxon:4100" /clone="nknhs1s"	callus tissue and root tissue"  (lab_host="DH10B-TonA"  /lab_host="DH10B-TonA"  /lab_host="NMicotiana benthamiana mixed tissue cDNA library normalised (1) 10 meth	/note="Vector: pCMVSports.liste 1: EcoR1; Site 2: Not1; supplier: RNA was isolated from Nicotiana benthamiana tissues that include callus, roots from liquid culture grown plants, heat-stressed leaves (38 C, 3 hr and 6 hr), cold-stressed leaves (5 C 3 hr, 6hr), and pathogen

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CGTGCTTTACGCTATCGCCGCTTCCCAGCGCATCGCCTTCTATCGCCTTCTTGA 1168
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EST704513 Nicotiana benthamiana mixed tissue cDNA library, normalized, full-length Nicotiana benthamiana cDNA clone NBMC477 5'
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1 (bases 1 to 933)
Buell, C.R., Hart, A. Zismann, V., Karamycheva, S.A., Day, B., Generation of EST sequences from Nicotiana benthamiana
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/clone lib="NH10B-TonA"
library, normalized, full-length"
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Other ESTS: EST754514
Other Ests: EST754514
Contact: Robin Buell
The Institute for Genomic Research
9712 Medical Center Dr. Rockville, MD 20850, USA
Email: potato-array@tigr.org
Email: potato-array@tigr.org
Institute via http://genome.arizona.edu/orders/
Seg primer: ATT TAG GTG ACA CTA TAG.
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library, normalized, full-length"
/note="Wector: pCNVSport6.1; Site 1: ECORI; Site 2: NotI;
/note="Wector: pCNVSport6.1; Site 1: ECORI; Site 2: NotI;
supplier: RNA was isolated from Nicotiana benthamiana
tissues that include callus, roots from liquid culture
grown plants, heat-etressed leaves (38 C, 3 hr and 6 hr),
cold-stressed leaves (5C 3 hr, 6hr), and pathogen
challenged leaves (5C 3 hr, 6hr), and pathogen
challenged leaves (Fecudomonas syringae pv tomato 12 hr;
Xanthomonas campestris pv campestris 12 hr, 18hr;
Pseudomonas syringae pv phaseolicola 18hr, and Kanthomonas
campestris pv vesicatoria 18hr). RNA was isolated from
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Bukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Bukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;

asterids; lamiids; Solanales; Solanaceae; Nicotiana.

1 (bases 1 to 910)

Staskawicz, B. (Jin,H. and Baker,B.

Generation of EST sequences from Nicotiana benthamiana

In Unpublished (2003)

Other ESTS: ESTS50653

Contact: Robin Buell

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Email: potato-array@tigr.org

Clones can be requested from the University of Arizona Genomics

Institute via http://genome.arizona.edu/orders/

Seq primer: ATT TAG GTG ACA CTA TAG.
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               normalized, full-length Nicotiana benthamiana cDNA clone NBMBC75 5'
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/note="Vector: pCMVSport6.1; Site 1: EcoRI; Site 2: NotI; supplier: RNA was isolated from Nicotiana benthamiana tissues that include callue, roots from liquid culture grown plants, heart-stressed leaves (38 C, 3 hr and 6 hr), cold-stressed leaves (5 C 3 hr, 6hr), and pathogen challenged leaves (Feeudomonas syringae pv tomato 12 hr; Xanthomonas campestris pv campestris 12 hr, 18hr; Pseudomonas syringae pv phaseolicola 18hr, and Xanthomonas campestris pv vesicatoria 18hr). RNA was isolated from these tissues and pooled in approximately equal molar
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CK256977 936 bp mRNA linear EST 30-JUL-2004 EST740614 potato callus cDNA library, normalized and full-length Solanum tuberosum cDNA clone POCD170 5' end, mRNA sequence.
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                                                                                                                                                                                                                                                                            Contact: Robin Buell
The Institute for Genomic Research
9712 Medical Center Dr. Rockville, MD 20850, USA
Email: potato-arrayofigr.org
Clones can be requested from the University of Arizona Genomics
Institute via http://genome.arizona.edu/orders/
Seq primer: ATT TAG GTG ACA CTA TAG.
Location/Qualifiers
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             /clone_lib="potato callus cDNA library, normalized and
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             /note="Vector: pCMVSport6.1; Site 1: EcoRI; Site 2: No supplier: RNA was isolated from Solanum tuberosum var. Kennebec callus tissue grown on solid media."
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Lases I to 938,
Buell, C.R., Hart, A., Zismann, V., Karamycheva, S.A. and Baker, Unpublished (203)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Length 936;
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100.0%; Pred. No. 3.9e-174;
tive 0; Mismatches 0;
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/organism="Solanum tuberosum"
                                                                                                                                                                                                                                                                                                                                                                                                                                         /mol_type="mRNA"
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/db_xref="taxon:4113"
/clone="POCD170"
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/lab_host="DH10B-TonA"
                                                                                                                                Solanum tuberosum (potato)
Solanum tuberosum
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Xanthomonas campestris pv campestris 12 hr, 18hr, Pseudomonas syringae pv phaseolicola 18hr, and Xanthomonas campestris pv vesicatoria 18hr). RNA was isolated from these tissues and pooled in approximately equal molar

1108 cgrectrracecrariceccecracteccarrecearceccrrcrarececrrrraca 1168 1048 919 736 964 868 496 928 556 919 376 436 628 688 cgacerrercacreaaecesaaaesaacrescrarreseseaaerescases 316 748 508 136 94 TCTCCTGTCATCTCACCTTGCTCCTGCCGAGAAAGTATCCATCATGGCTGATGCAATGCG GCGGCTGCATACGCTTGATCCGGCTACCTGCCCATTCGACCCACCAAGCGAAACATCGCAT CGAGCGAGCACGTACTCGGATGGGAAGCCGGTCTTGTCGATCAGGATGATCTGGACGAAGA CGAGGATCTCGTCGTCACCCATGCCTGCTTGCCGAATATCATGGTGGAAAATGG CCGCTTTTCTCGATTCATCGACTGTGCCCGCTGGGTGTGGCGGACCGCTATCAGGACAT AGCGTTGGCTACCCGTGATATTGCTGAAGAGCTTGGCGGCGAATGGGCTGACCGCTTCCT 737 CGIGCTITACGGIAICGCCGCICCCGAITCGCAGCGCAICGCCITCTAICGCCCTTCTTGA GCGGCTGCATACGCTTCGGCTACCTGCCCATTCGACCACCAAGCGAAACATCGCAT GGACGAGGCAGCGAGCTATCGTGGCTGGCCACGAGGGGGTTCCTTGCGCAGCTGTGCT CGACGTTGTCACTGAAGCGGGAAGGGACTGGCTGCTATTGGGCGAAGTGCCGGGGCAAGA GATTGAACAAGATGGATTGCACGCAGGTTCTCCGGCCGCTTGGGTGGAGAGGCTATTCGG 17 GALTGAACAAGATGGATGCACGCAGGTTCTCCGGCCGCTTGGGTGGAGAGGCTATTCGG CTATGACTGGGCACAACAGACAATCGGCTGCTCTGATGCCGCCGTGTTCCGGCTGTCAGC Gaps

CK283361 BST702-AUG-2004 BST746083 Nicotiana benthamiana mixed tissue cDNA library, normalized, full-length Nicotiana benthamiana cDNA clone NBMAG50 5'

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CK288185.1 GI:39865462
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/lab_hot="maltotic and biotic stress-treated leaves,
/lab_hot="maltotic and root tissue"
/lab_hot="maltotic form"
/clone_lib="wicotic manalized, full-length"
/clone_lib="wicotic: pCMVSport6.1; Site 1: BCORI; Site 2: NotI;
/note="weetor: pCMVSport6.1; Site 1: BCORI; Site 2: NotI;
supplier: RNA was isolated from Nicotiana benthamiana
tissues that include callus, roots from liquid culture
grown plants, heat-stressed leaves (38 C, 3 hr and 6 hr),
cold-stressed leaves (5 C shr, and pathogen
challenged leaves (5 C shr, and pathogen
xanthomonas campestris pv campestris 12 hr, 18hr;
Pseudomonas syringae pv phaseolicola 18hr; and Xanthomonas
campestris pv vesicatoria 18hr). RNA was isolated from
these tissues and pooled in approximately equal molar
                                                                         Nicotiana benthamiana
Bukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
asterids; laminds; Solanales; Solanaceae; Nicotiana.
1 (bases 1 to 954)
Buell, C.R., Hart, A., Zismann, V., Karamycheva, S.A., Day, B.,
Staskawicz, B., Jin, H. and Baker, B.
Generation of EST sequences from Nicotiana benthamiana
Other ESTs: EST746084
Contact: Robin Buell
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                                                                                                                                                                                                                                                          The Institute for Genomic Research 9712 Medical Center Dr, Rockville, MD 20850, USA Bmail: potato-array@tigr.org Clones can be requested from the University of Arizona Genomics Institute via http://genome.arizona.edu/orders/ . Seq primer: ATT TAG GTG ACA CTA TAG.
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100.0%; Pred. No. 3.9e-174;
iive 0; Mismatches 0; Indels
                                                                                                                                                                                                                                                                                                                                                                                               organism="Nicotiana benthamiana"
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/db_xref="taxon:4100"
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                                CK283361.1 GI:39855898
                                                               Nicotiana benthamiana
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Micotiana benthamiana
Bukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
asterids; lamiids; Solanales; Solanaceae; Nicotiana.
1 (bases 1 to 811)
Buelli,C.R., Hart,A., Zismann,V., Karamycheva,S.A., Day,B.,
Generation of EST sequences from Nicotiana benthamiana
Unpublished (2003)
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/lab_host='DhlBB-TonA"
/clone lib="Nicotiana benthamiana mixed tissue cDNA
library, normalized, full-length*
/note="Vector: pGMVSport6.1; Site 1: EcoR1; Site 2: Not1; supplier: RNA was isolated from Nicotiana benthamiana
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                                 374 GCGGCTGCATACGCTTGATCCGGCTACCTGCCCATTCGACCACCAACGAACATCGCCAT
                                                                                                     809 CGAGCGAGCACGTACTCGGATGGAAGCCGGTCTTGTCGATCAGGATGATCTGGACGAAGA
                                                                                                                                                     434 CGAGCGAGCACGTACTCCGGATGGAAGCCGGTCTTGTCGATCAGGATGATCTGGACGAAGA
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9712 Medical Center Dr, Rockville, MD 20850, USA
Email: potato-array@tigr.org
Clones can be requested from the University of Arizona Genomics
Institute via http://genome.arizona.edu/orders/
Seg primer: ATT TAG GTG ACA CTA TAG.
Location/Qualifiers
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/mol_type="mRNA"
/db_xref="taxon:4100"
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1045
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tissues that include callus, roots from liquid culture grown plants, heat-stressed leaves (38 C, 3 hr and 6 hr), cold-stressed leaves (5 C 3 hr, 6hr), and pathogen challenged leaves (Fseudomos syringae py tomato 12 hr; Xanthomonas campestris py campestris 12 hr, 18hr; Pseudomonas syringae py phaseolicola 18hr, and Xanthomonas campestris py vesicatoria 18hr). RNA was isolated from these tissues and pooled in approximately equal molar amounts."
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                                                                                                                                                           9.7%; Score 778.8; DB 7;
llarity 99.7%; Pred. No. 1.3e-170;
Conservative 0; Mismatches 2;
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Matches 780; Conserv
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/lab host="BH10B-TonA"
/lab host="BH10B-TonA"
/clome lib="Wicotic hours liberary, normalized, full-length"
/lonce="Weetor: pCWNSport6.1; Site_1: EcoRI; Site_2: NotI;
/note="Weetor: pCWNSport6.1; Site_1: EcoRI; Site_2: NotI;
supplier: RNA was isolated from Nicotiana benthamiana
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grown plants, heat-stressed leaves (8s C, 3 hr, old-stressed leaves (5 C 3 hr, 6hr), and pathogen
cold-stressed leaves (Fseudomonas syringae pv tomato 12 hr;
Xanthomonas campestris pv campestris 12 hr, 18hr;
Pseudomonas syringae pv phaseolicola 18hr, and Xanthomonas
campestris pv vesicatoria 18hr). RNA was isolated from
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      CK289711 BST 02-AUG-2004 EST751433 Nicotiana benthamiana mixed tissue cDNA library, normalized, full-length Nicotiana benthamiana cDNA clone NBMBI49 5'
                                                                                                                                                                              Eukaryotta; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Eperatophyta; Magnoliophyta; eudicotyledons; core eudicots; asterids; lamiids; Solanales; Solanaceae; Nicotiana.

1 (Dases 1 to 878)

Buell, C.R., Hart, A., Zismann, V., Karamycheva, S.A., Day, B., Staskawicz, B., Jin, H. and Baker, B. Generation of EST sequences from Nicotiana benthamiana
Unpublished (2003)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    9
                                                                                                                                                                                                                                                                                                                                  Contact: Robin Buell
The Institute for Genomic Research
9712 Medical Center Dr, Rockville, MD 20850, USA
Bmail: pocato-array@tigr.org
Clones can be requested from the University of Arizona Genomics
Institute via http://genome.arizona.edu/orders/
Seq primer: ATT TAG GTG ACA CTA TAG.
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Pred. No. 9.3e-126;
0; Mismatches 0;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        /organism="Nicotiana benthamiana"
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       /mol_type="mRNA"
/db_xref="taxon:4100"
/clone="NBMBI49"
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                                                                                                                                                                                 TCACCTIGCTCCTGCCGAGAAAGTAICCATCATGGCTGATGCGGCGGCTGCATAC 760
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Yu,Y., Zhu,H., Boyd,C.A., Gaudette,B., Gayle,A., Kingsbury,R., Phillips,K., Sasinowski,M, Wing,R.A. and Dean,R.A.
A BAC End Sequencing Framework to Sequence the Magnaporthe grisea
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Sordariomycetes incertae sedis; Magnaporthaceae; Magnaporthe.
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Magnaporthe grisea
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Clemson University
100 Vordan Hall, Clemson University, Clemson,
Fax: 864 656 5737
                                                            13;
                    DB 8;
                    Score 577.4; DB 8;
Pred. No. 1.6e-123;
                                                      0; Mismatches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Seq primer: TAATACGACTCACTATAGGG
Class: BAC ends
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GSS.
           Query Match 7.2%;
Best Local Similarity 97.8%;
Matches 584; Conservative
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/lab_host="E. coli DH10B"
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/note="Vector: pBACWICH; Site_1: HindIII; Site 2: HindIII;
Rice blast is one of the most devestating fungal diseases
of rice world wide. It is a filamentous ascomycete with
a haploid genome (n=7) of approximately 40 Mbp. Rice
blast is an important model fungal pathogen for studying
numerous aspects of the fungal-host interaction. In
order to facilitate genome wide analysis, a BAC library
containing 921c clones with an average insert size of 130
kbp was constructed. This library represents greater
than 25X genome coverage. High density colony filters
are available upon request."
                                                                      TGCCTGCTTGCCGAATATCATGGTGGAAAATGGCCGCTTTTCTGGATTCATCGACTGTGG 1015
                                                                                                                                                                                                                  AGAGCTIGGCGGCGAATGGGCTGACCGCTTCCTCGTGCTTTACGGTATCGCCGCTCCCGA 1135
                                                                                                                                            1016 CCGGCTGGGTGTGGCGGACCGCTATCAGGACATAGCGTTGGCTACCCGTGATATTGCTGA 1075
                                                                                                                                                                                                                                                                                                                                                                                                                    AQ361914 1789 bp DNA linear GSS 03-FBB-1999 mgxb0005K01f CUGI Rice Blast BAC Library Magnaporthe grisea genomic
                                               360
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           955
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A BAC End Sequencing Framework to Sequence the Magnaporthe grisea
                        GTTCGCCAGGCTCAAGGCGCGCGATGCCCGACGAGGATCTCGTCGTGACCCATGGCGA
                                                                                                                                                                  Pezizomycotina; Sordariomycetes;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Magnaporthe grisea

Eukaryota; Fungi; Ascomycota; Pezizomycotina; Sordariomycetes
Sordariomycetes incertae sedis; Magnaporthaceae; Magnaporthe.

1 (bases 1 to 789)
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AL Unpublished (1998)
Contact: Dean RA
Contact: Dean RA
Contact: Dean RA
Clemson University
100 Jordan Hall, Clemson Universiy, Clemson, SC 29634
Tel: 864 656 5737
Fax: 864 656 4293
Email: rdean@clemson.edu
Seq primer: TAATACGACTATAGGG
Class: BAC ends
High quality sequence start: 41
High quality sequence stop: 392.
Location/Qualifiers

1 789
                                                                                                                                                                                                                                                                                        TTCGCAGCGCATCGCCTTCTATCGCCTTCTTGACGAGTTCTTCTGAG 1182
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TITLE

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High quality sequence stop: 187

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976 TGGTGGAAAATGGCCGCTTTTCTGGATTCATCGACTGTGGGCGGGTGTGGGCGGACC 1035
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Fax: 405 325 7762
Email: broecou.edu
Contact Dr.Rouf Mian (rmian@noble.org) regarding clone availability
Seq primer: M13 reverse primer
High quality sequence stop: 470.
                                                                                                                     1 (bases 1 to 549)
Zhang, Y., Zwonitzer, J.C., Chekhovskiy, K., May, G.D. and Mian, M.A.R.
A functional genemics approach for identification of heat tolerance genes in tall fescue
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            246 CGAAACATCGCATCGAGCGAGCACGTACTCGGATGGAAGCCGGTCTTGTCGATCAGGATG 305
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1; BD/Clontech PCR-select cDNA subtraction library"
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                                                            Schedonorus arundinaceus
Bukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Pooideae; Poeae; Schedonorus.
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Department of Chemistry and Biochemistry
Advanced Center for Genome Technology, University of Oklahoma
620 Parzington Oval, Norman, OK 73019, USA
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                                        Schedonorus arundinaceus (Festuca arundinacea)
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/organism="Schedonorus arundinaceus"
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            GI:43400943
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                                                                                                                     /tissue_type="Protoplasts"
/tissue_type="Protoplasts"
/lab host="E. coli Bilast BAC Library"
/clone_lib="CUGI Rice Blast BAC Library"
/clone_lib="VCGI Rice Blast BAC Library"
/clone_lib="VCGI Rice Blast BAC Library"
/clone_station of the most devestating fungal diseases
Rice blast is one of the most devestating fungal diseases
of rice world wide. It is a filamentous ascomycete with
a haploid genome (n=7) of approximately 40 Mbp. Rice
blast is an important model fungal pathogen for studying
numerous aspects of the fungal-host interaction. In
order to facilitate genome wide analysis, a BAC library
containing 9216 clones with an average insert size of 130
kbp was constructed. This library represents greater
than 25X genome coverage. High density colony filters
are available upon request."
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Matches 564; Conservative 0; Mismatches 7; Indels 0
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                                    organism="Magnaporthe grisea"
(mol_type="genomic DNA"
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AQ44775 509 bp DNA linear GSS 08-APR-1999 mgxb0011E13f CUGI Rice Blast BAC Library Magnaporthe grisea genomic clone mgxb0011E13f, genomic survey sequence.
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/note="Vector: PBACWICH; Site_1: HindIII; Site_2: HindIII;
Rice blast is one of the most devestating fungal diseases of rice world wide. It is a filamentous ascomycete with
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Yu,Y., Zhu,H., Boyd,C.A., Gaudette,B., Gayle,A., Kingsbury,R., Phillips,K., Sasinowski,M, Wing,R.A. and Dean,R.A.
A BAC End Sequencing Framework to Sequence the Magnaporthe grisea Genome
                                                                         GACGITGTCACTGAAGCGGGAAGGGACTGCTATTGGGCGAAGTGCCGGGGCAGGAT
                                                                                                                                                CTCCTGTCATCTCACCTTGCTCCTGCCGAGAAGTATCCATCATGGCTGATGCAATGCGG
                                                                                              CGGCTGCATACGCTTGATCCGGCTACCTGCCCATTCGACCACCAAGCGAAACATCGCATC
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                                                                                                                                                                                                                                                                                                                     CATCAGGGGCTCGCCAGCCGAACTGTTCGCCAGGCTCAAAGGCGCGCATGCCCGACGGC
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Contact: Dean RA
Clemson University Genomics Institute
clemson University
100 Jordan Hall, Clemson Universiy, Clemson, SC 29634
Tel: 864 656 5737
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Magnaporthe grisea (anamorph: Pyricularia grisea)
Magnaporthe grisea
Eukaryota; Fungi; Ascomycota; Pezizomycotina; Sor
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/strain="70-15"
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Email: rdean@clemson.edu
Seq primer: TAATACGACTCACTATAGGG
Class: BAC ends
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Location/Qualifiers
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/tissue_type="Protoplasts"
/lab host="E. coli DH10B"
/clone_lib="CUGI Rice Blast BAC Library"
/note="Vector: PBACWICH; Site_l: HindII; Site_2: HindIII;
Rice blast is one of the most devestating fungal diseases of rice world wide. It is a filamentous ascomycete with
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            a haploid genome (n=7) of approximately 40 Mbp. Rice
blast is an important model fungal pathogen for studying
numerous aspects of the fungal host interaction. In
order to facilitate genome wide analysis, a BAC library
containing 9216 close with an average insert size of 130
kbp was constructed. This library represents greater
than 25X genome coverage. High density colony filters
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Yu,Y., Zhu,H., Boyd,C.A., Gaudette,B., Gayle,A., Kingsbury,R.,
Phillips,K., Sasinowski,M, Wing,R.A. and Dean,R.A.
A BAC End Sequencing Framework to Sequence the Magnaporthe grisea
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                                                                            Magnaporthe grisea (anamorph: Pyricularia grisea)
Magnaporthe grisea
Eukaryota; Fungi; Ascomycota; Pezizomycotina; Sordariomycetes;
Sordariomycetes incertae sedis; Magnaporthaceae; Magnaporthe.
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Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson Universiy, Clemson, SC 29634
Tel: 864 656 5737
Fax: 864 656 4293
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Length 561;
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Pred. No. 6.4e-109;
0; Mismatches 2;
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High quality sequence stop: 326.
Location/Qualifiers
1..561
/organism="Magnaporthe grisea"
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/strain="70-15"
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Seg primer: TAATACGACTCACTATAGGG
Class: BAC ends
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Best Local Similarity 99.6%;
Matches 516; Conservative
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a haploid genome (n=7) of approximately 40 Mpp. Rice blast is an important model fungal pathogen for studying numerous aspects of the fungal-host interaction. In order to facilitate genome wide analysis, a BAC library containing 9216 clones with an average insert size of 130 kpp was constructed. This library represents greater khan 25x genome coverage. High density colony filters are available upon request."
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ö Gaps .. Length 509; Indels Query Match 6.4%; Score 508; DB 8; Le Best Local Similarity 100.0%; Pred. No. 2.4e-107; Matches 508; Conservative 0; Mismatches 0; ORIGIN

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